

NIST SPECIAL PUBLICATION 260-131

U.S. DEPARTMENT OF COMMERCE/Technology Administration National Institute of Standards and Technology

Standard Reference Materials:

The Certification of 100 mm Diameter Silicon Resistivity SRMs 2541 through 2547 Using Dual-Configuration Four-Point Probe Measurements

J. R. Ehrstein and M. C. Croarkin

he National Institute of Standards and Technology was established in 1988 by Congress to "assist industry in the development of technology . . . needed to improve product quality, to modernize manufacturing processes, to ensure product reliability . . . and to facilitate rapid commercialization . . . of products based on new scientific discoveries."

NIST, originally founded as the National Bureau of Standards in 1901, works to strengthen U.S. industry's competitiveness; advance science and engineering; and improve public health, safety, and the environment. One of the agency's basic functions is to develop, maintain, and retain custody of the national standards of measurement, and provide the means and methods for comparing standards used in science, engineering, manufacturing, commerce, industry, and education with the standards adopted or recognized by the Federal Government.

As an agency of the U.S. Commerce Department's Technology Administration, NIST conducts basic and applied research in the physical sciences and engineering, and develops measurement techniques, test methods, standards, and related services. The Institute does generic and precompetitive work on new and advanced technologies. NIST's research facilities are located at Gaithersburg, MD 20899, and at Boulder, CO 80303. Major technical operating units and their principal activities are listed below. For more information contact the Publications and Program Inquiries Desk, 301-975-3058.

Office of the Director

- National Quality Program
- · International and Academic Affairs

Technology Services

- Standards Services
- Technology Partnerships
- Measurement Services
- Technology Innovation
- · Information Services

Advanced Technology Program

- Economic Assessment
- Information Technology and Applications
- Chemical and Biomedical Technology
- · Materials and Manufacturing Technology
- · Electronics and Photonics Technology

Manufacturing Extension Partnership Program

- · Regional Programs
- · National Programs
- · Program Development

Electronics and Electrical Engineering Laboratory

- Microelectronics
- Law Enforcement Standards
- Electricity
- Semiconductor Electronics
- Electromagnetic Fields¹
- Electromagnetic Technology¹
- Optoelectronics¹

Chemical Science and Technology Laboratory

- Biotechnology
- Physical and Chemical Properties²
- Analytical Chemistry
- Process Measurements
- Surface and Microanalysis Science

Physics Laboratory

- · Electron and Optical Physics
- Atomic Physics
- · Optical Technology
- Ionizing Radiation
- Time and Frequency¹
- · Quantum Physics1

Materials Science and Engineering Laboratory

- · Intelligent Processing of Materials
- Ceramics
- Materials Reliability
- Polymers
- Metallurgy
- · NIST Center for Neutron Research

Manufacturing Engineering Laboratory

- Precision Engineering
- · Automated Production Technology
- · Intelligent Systems
- · Fabrication Technology
- · Manufacturing Systems Integration

Building and Fire Research Laboratory

- Structures
- · Building Materials
- Building Environment
- · Fire Safety Engineering
- Fire Science

Information Technology Laboratory

- Mathematical and Computational Sciences²
- Advanced Network Technologies
- · Computer Security
- · Information Access and User Interfaces
- High Performance Systems and Services
- Distributed Computing and Information Services
- Software Diagnostics and Conformance Testing

¹At Boulder, CO 80303.

²Some elements at Boulder, CO.

NIST Special Publication 260-131

Standard Reference Materials:

The Certification of 100 mm Diameter Silicon Resistivity SRMs 2541 through 2547 Using Dual-Configuration Four-Point Probe Measurements

J. R. Ehrstein

Semiconductor Electronics Division Electronics and Electrical Engineering Laboratory

M. C. Croarkin

Statistical Engineering Division Information Technology Laboratory

National Institute of Standards and Technology Gaithersburg, MD 20899-0001



U.S. DEPARTMENT OF COMMERCE, William M. Daley, Secretary TECHNOLOGY ADMINISTRATION, Gary R. Bachula, Acting Under Secretary for Technology NATIONAL INSTITUTE OF STANDARDS AND TECHNOLOGY, Robert E. Hebner, Acting Director

Issued August 1997

National Institute of Standards and Technology Special Publication 260–131 Natl. Inst. Stand. Technol. Spec. Publ. 260–131, 98 pages (Aug. 1997) CODEN: NSPUE2

U.S. GOVERNMENT PRINTING OFFICE WASHINGTON: 1997

Foreword

Standard Reference Materials (SRMs) as defined by the National Institute of Standards and Technology (NIST) are well-characterized materials, produced in quantity and certified for one or more physical or chemical properties. They are used to assure the accuracy and compatibility of measurements throughout the Nation. SRMs are widely used as primary standards in many diverse fields in science, industry, and technology, both within the United States and throughout the world. They are also used extensively in the fields of environmental and clinical analysis. In many applications, traceability of quality control and measurement processes to the national measurement system is carried out through the mechanism and use of SRMs. For many of the Nation's scientists and technologists, it is therefore of more than passing interest to know the details of the measurements made at NIST in arriving at the certified values of the SRMs produced. The NIST Special Publication 260 Series is a series of papers reserved for this purpose.

The 260 Series is dedicated to the dissemination of information on different phases of the preparation, measurement, certification, and use of NIST SRMs. In general, much more detail will be found in these papers than is generally allowed, or desirable, in scientific journal articles. This enables the user to assess the validity and accuracy of the measurement processes employed, to judge the statistical analysis, and to learn details of techniques and methods utilized for work entailing greatest care and accuracy. These papers also should provide sufficient additional information so SRMs can be utilized in new applications in diverse fields not foreseen at the time the SRM was originally issued.

Inquiries concerning the technical content of this paper should be directed to the author(s). Other questions concerned with the availability, delivery, price, and so forth, will receive prompt attention from:

Standard Reference Materials Program Bldg. 202, Rm. 204
National Institute of Standards and Technology Gaithersburg, MD 20899
Telephone: (301) 975-6776
FAX: (301) 948-3730

e-mail: srminfo@nist.gov, or www:http://ts.nist.gov/srm

Thomas E. Gills, Chief Standard Reference Materials Program

OTHER NIST PUBLICATIONS IN THIS SERIES

- Trahey, N.M., ed., NIST Standard Reference Materials Catalog 1995-96, NIST Spec. Publ. 260 (1995 Ed.). PB95-232518/AS
- Michaelis, R.E., and Wyman, L.L., Standard Reference Materials: Preparation of White Cast Iron Spectrochemical Standards, NBS Misc. Publ. 260-1 (June 1964). COM74-11061**
- Michaelis, R.E., Wyman, L.L., and Flitsch, R., Standard Reference Materials: Preparation of NBS Copper-Base Spectrochemical Standards, NBS Misc. Publ. 260-2 (October 1964). COM74-11063**
- Michaelis, R.E., Yakowitz, H., and Moore, G.A., Standard Reference Materials: Metallographic Characterization of an NBS Spectrometric Low-Alloy Steel Standard, NBS Misc. Publ. 260-3 (October 1964). COM74-11060**
- Hague, J.L., Mears, T.W., and Michaelis, R.E., Standard Reference Materials: Sources of Information, Publ. 260-4 (February 1965). COM74-11059**
- Alvarez, R., and Flitsch, R., Standard Reference Materials: Accuracy of Solution X-Ray Spectrometric Analysis of Copper-Base Alloys, NBS Misc. Publ. 260-5 (February 1965). PB168068**
- Shultz, J.I., Standard Reference Materials: Methods for the Chemical Analysis of White Cast Iron Standards, NBS Misc. Publ. 260-6 (July 1965). COM74-11068**
- Bell, R.K., Standard Reference Materials: Methods for the Chemical Analysis of NBS Copper-Base Spectrochemical Standards, NBS Misc. Publ. 260-7 (October 1965). COM74-11067**
- Richmond, M.S., Standard Reference Materials: Analysis of Uranium Concentrates at the National Bureau of Standards, NBS Misc. Publ. 260-8 (December 1965). COM74-11066**
- Anspach, S.C., Cavallo, L.M., Garfinkel, S.B., et al., Standard Reference Materials: Half Lives of Materials Used in the Preparation of Standard Reference Materials of Nineteen Radioactive Nuclides Issued by the National Bureau of Standards, NBS Misc. Publ. 260-9 (November 1965). COM74-11065**

- Yakowitz, H., Vieth, D.L., Heinrich, K.F.J., et al., Standard Reference Materials: Homogeneity Characterization of NBS Spectrometric Standards II: Cartridge Brass and Low-Alloy Steel, NBS Misc. Publ. 260-10 (December 1965). COM74-11064**
- Napolitano, A., and Hawkins, E.G., Standard Reference Materials: Viscosity of Standard Lead-Silica Glass, NBS Misc. Publ. 260-11** (November 1966).
- Yakowitz, H., Vieth, D.L., and Michaelis, R.E., Standard Reference Materials: Homogeneity Characterization of NBS Spectrometric Standards III: White Cast Iron and Stainless Steel Powder Compact, NBS Misc. Publ. 260-12 (September 1966).
- Spijkerman, J.J., Snediker, D.K., Ruegg, F.C., et al., Standard Reference Materials: Mossbauer Spectroscopy Standard for the Chemical Shift of Iron Compounds, NBS Misc. Publ. 260-13** (July 1967).
- Menis, O., and Sterling, J.T., Standard Reference Materials: Determination of Oxygen in Ferrous Materials (SRMs 1090, 1091, 1092), NBS Misc. Publ. 260-14** (September 1966).
- Passaglia, E. and Shouse, P.J., Standard Reference Materials: Recommended Method of Use of Standard Light-Sensitive Paper for Calibrating Carbon Arcs Used in Testing Testiles for Colorfastness to Light, NBS Spec. Publ. 260-15 (July 1967). Superseded by SP 260-41.
- Yakowitz, H., Michaelis, R.E., and Vieth, D.L., Standard Reference Materials: Homogeneity Characterization of NBS Spectrometric Standards IV: Preparation and Microprobe Characterization of W-20% Mo Alloy Fabricated by Powder Metallurgical Methods, NBS Spec. Publ. 260-16 (January 1969). COM74-11062**
- Catanzaro, E.J., Champion, C.E., Garner, E.L., et al., Standard Reference Materials: Boric Acid; Isotopic, and Assay Standard Reference Materials, NBS Spec. Publ. 260-17 (February 1970). PB189457**

- Geller, S.B., Mantek, P.A., and Cleveland, N.G., Calibration of NBS Secondary Standards Magnetic Tape Computer Amplitude Reference Amplitude Measurement "Process A," NBS Spec. Publ. 260-18 (November 1969). Superseded by SP 260-29.
- Paule, R.C., and Mandel, J., Standard Reference Materials: Analysis of Interlaboratory Measurements on the Vapor Pressure of Gold (Certification of SRM 745). NBS Spec. Publ. 260-19 (January 1970). PB190071**
- 260-20: Unassigned
- Paule, R.C., and Mandel, J., Standard Reference Materials: Analysis of Interlaboratory Measurements on the Vapor Pressures of Cadmium and Silver, NBS Spec. Publ. 260-21 (January 1971). COM74-11359**
- Yakowitz, H., Fiori, C.E., and Michaelis, R.E., Standard Reference Materials: Homogeneity Characterization of Fe-3 Si Alloy, NBS Spec. Publ. 260-22 (February 1971). COM74-11357**
- Napolitano, A., and Hawkins, E.G., Standard Reference Materials: Viscosity of a Standard Borosilicate Glass, NBS Spec. Publ. 260-23 (December 1970). COM71-00157**
- Sappenfield, K.M., Marinenko, G., and Hague, J.L., Standard Reference Materials: Comparison of Redox Standards, NBS Spec. Publ. 260-24 (January 1972). COM72-50058***
- Hicho, G.E., Yakowitz, H., Rasberry, S.D., et al., Standard Reference Materials: A Standard Reference Material Containing Nominally Four Percent Austenite, NBS Spec. Publ. 260-25 (February 1971). COM74-11356**
- Martin, J.F., Standard Reference Materials: NBS-U.S. Steel Corp. Joint Program for Determining Oxygen and Nitrogen in Steel, NBS Spec. Publ. 260-26 (Γebruary 1971). PB 81176620**
- Garner, E.L., Machlan, L.A., and Shields, W.R., Standard Reference Materials: Uranium Isotopic Standard Reference Materials, NBS Spec. Publ. 260-27 (April 1971). COM74-11358**

- Heinrich, K.F.J., Myklebust, R.L., Rasberry, S.D., et al., Standard Reference Materials: Preparation and Evaluation of SRMs 481 and 482 Gold-Silver and Gold-Copper Alloys for Microanalysis, NBS Spec. Publ. 260-28 (August 1971). COM71-50365**
- Geller, S.B., Standard Reference Materials:
 Calibration of NBS Secondary Standard
 Magnetic Tape (Computer Amplitude Reference)
 Using the Reference Tape Amplitude
 Measurement "Process A-Model 2," NBS Spec.
 Publ. 260-29 (June 1971). COM71-50282**
 Supersedes Measurement System in SP 260-18.
- Gorozhanina, R.S., Freedman, A.Y., and Shaievitch, A.B., (translated by M.C. Selby), Standard Reference Materials: Standard Samples Issued in the USSR (A Translation from the Russian), NBS Spec. Publ. 260-30 (June 1971). COM71-50283**
- Hust, J.G., and Sparks, L.L., Standard Reference Materials: Thermal Conductivity of Electrolytic Iron SRM 734 from 4 to 300 K, NBS Spec. Publ. 260-31 (November 1971). COM71-50563**
- Mavrodineanu, R., and Lazar, J.W., Standard Reference Materials: Standard Quartz Cuvettes for High Accuracy Spectrophotometry, NBS Spec. Publ. 260-32 (December 1973).COM74-50018***
- Wagner, H.L., Standard Reference Materials: Comparison of Original and Supplemental SRM 705, Narrow Molecular Weight Distribution Polystyrene, NBS Spec. Publ. 260-33 (May 1972). COM72-50526**
- Sparks, L.L., and Hust, J.G., Standard Reference Material: Thermoelectric Voltage of Silver-28 Atomic Percent Gold Thermocouple Wire, SRM 733, Verses Common Thermocouple Materials (Between Liquid Helium and Ice Fixed Points), NBS Spec. Publ. 260-34 (April 1972). COM72-50371**
- Sparks, L.L., and Hust, J.G., Standard Reference Materials: Thermal Conductivity of Austenitic Stainless Steel, SRM 735 from 5 to 280 K, NBS Spec. Publ. 260-35 (April 1972). COM72-50368**

- Cali, J.P., Mandel, J., Moore, L.J., et al., Standard Reference Materials: A Reference Method for the Determination of Calcium in Serum NBS SRM 915, NBS Spec. Publ. 260-36 (May 1972). COM72-50527**
- Shultz, J.I., Bell, R.K., Rains, T.C., et al., Standard Reference Materials: Methods of Analysis of NBS Clay Standards, NBS Spec. Publ. 260-37 (June 1972). COM72-50692**
- Richard, J.C., and Hsia, J.J., Standard Reference Materials: Preparation and Calibration of Standards of Spectral Specular Reflectance, NBS Spec. Publ. 260-38 (May 1972). COM72-50528***
- Clark, A.F., Denson, V.A., Hust, J.G., et al., Standard Reference Materials: The Eddy Current Decay Method for Resistivity Characterization of High-Purity Metals, NBS Spec. Publ. 260-39 (May 1972). COM72-50529**
- McAdie, H.G., Garn, P.D., and Menis, O., Standard Reference Materials: Selection of Differential Thermal Analysis Temperature Standards Through a Cooperative Study (SRMs 758, 759, 760), NBS Spec. Publ. 260-40 (August 1972) COM72-50776**
- Wood. L.A., and Shouse, P.J., Standard Reference Materials: Use of Standard Light-Sensitive Paper for Calibrating Carbon Arcs Used in Testing Textiles for Colorfastness to Light, NBS Spec. Publ. 260-41 (August 1972). COM72-50775**
- Wagner, H.L., and Verdier, P.H., eds., Standard Reference Materials: The Characterization of Linear Polyethylene, SRM 1475, NBS Spec. Publ. 260-42 (September 1972). COM72-50944**
- Yakowitz, H., Ruff, A.W., and Michaelis, R.E., Standard Reference Materials: Preparation and Homogeneity Characterization of an Austenitic Iron-Chromium-Nickel Alloy, NBS Spec. Publ. 260-43 (November 1972). COM73-50760**
- Schooley, J.F., Soulen, R.J., Jr., and Evans, G.A., Jr., Standard Reference Materials: Preparation and Use of Superconductive Fixed Point Devices, SRM 767, NBS Spec. Publ. 260-44 (December 1972). COM73-50037**

- Greifer, B., Maienthal, E.J., Rains, T.C., et al., Standard Reference Materials: Development of NBS SRM 1579 Powdered Lead-Based Paint, NBS Spec. Publ. 260-45 (March 1973). COM73-50226**
- Hust, J.G., and Giarratano, P.J., Standard Reference Materials: Thermal Conductivity and Electrical Resistivity Standard Reference Materials: Austenitic Stainless Steel, SRMs 735 and 798, from 4 to 1200 K, NBS Spec. Publ. 260-46 (March 1975). COM75-10339**
- Hust, J.G., Standard Reference Materials: Electrical Resistivity of Electrolytic Iron, SRM 797, and Austenitic Stainless Steel, SRM 798, from 5 to 280 K, NBS Spec. Publ. 260-47 (February 1974). COM74-50176**
- Mangum, B.W., and Wise, J.A., Standard Reference Materials: Description and Use of Precision Thermometers for the Clinical Laboratory, SRM 933 and SRM 934, NBS Spec. Publ. 260-48 (May 1974). Superseded by NIST Spec. Publ. 260-113. COM74-50533**
- Carpenter, B.S., and Reimer, G.M., Standard Reference Materials: Calibrated Glass Standards for Fission Track Use, NBS Spec. Publ. 260-49 (November 1974), COM74-51185**
- Hust, J.G., and Giarratano, P.J., Standard Reference Materials: Thermal Conductivity and Electrical Resistivity Standard Reference Materials: Electrolytic Iron, SRMs 734 and 797 from 4 to 1000 K, NBS Spec. Publ. 260-50 (June 1975). COM75-10698**
- Mavrodineanu, R., and Baldwin, J.R., Standard Reference Materials: Glass Filters As a SRM for Spectrophotometry-Selection, Preparation, Certification, and Use-SRM 930 NBS Spec. Publ. 260-51 (November 1975). COM75-10339**
- Hust, J.G., and Giarratano, P.J., Standard Reference Materials: Thermal Conductivity and Electrical Resistivity SRMs 730 and 799, from 4 to 3000 K, NBS Spec. Publ. 260-52 (September 1975). COM75-11193**
- Durst, R.A., Standard Reference Materials: Standardization of pH Measurements, NBS Spec. Publ. 260-53 (December 1978). Superseded by SP 260-53 Rev. 1988 Edition. PB88217427**

- Burke, R.W., and Mavrodineanu, R., Standard Reference Materials: Certification and Use of Acidic Potassium Dichromate Solutions as an Ultraviolet Absorbance Standard, NBS Spec. Publ. 260-54 (August 1977). PB272168**
- Ditmars, D.A., Cezairliyan, A., Ishihara, S., et al., Standard Reference Materials: Enthalpy and Heat Capacity; Molybdenum SRM 781, from 273 to 2800 K, NBS Spec. Publ. 260-55 (September 1977). PB272127**
- Powell, R.L., Sparks, L.L., and Hust, J.G., Standard Reference Materials: Standard Thermocouple Material, Pt-67: SRM 1967, NBS Spec. Publ. 260-56 (February 1978). PB277172**
- Cali, J.P., and Plebanski, T., Standard Reference Materials: Guide to United States Reference Materials, NBS Spec. Publ. 260-57 (February 1978). PB277173**
- Barnes, J.D., and Martin, G.M., Standard Reference Materials: Polyester Film for Oxygen Gas Transmission Measurements SRM 1470, NBS Spec. Publ. 260-58 (June 1979). PB297098**
- Chang, T., and Kahn, A.H., Standard Reference Materials: Electron Paramagnetic Resonance Intensity Standard: SRM 2601; Description and Use, NBS Spec. Publ. 260-59 (August 1978). PB292097**
- Velapoldi, R.A., Paule, R.C., Schaffer, R., et al., Standard Reference Materials: A Reference Method for the Determination of Sodium in Serum, NBS Spec. Publ. 260-60 (August 1978). PB286944**
- Verdier, P.H., and Wagner, H.L., Standard Reference Materials: The Characterization of Linear Polyethylene (SRMs 1482, 1483, 1484), NBS Spec. Publ. 260-61 (December 1978). PB289899**
- Soulen, R.J., and Dove, R.B., Standard Reference Materials: Temperature Reference Standard for Use Below 0.5 K (SRM 768), NBS Spec. Publ. 260-62 (April 1979). PB294245**
- Velapoldi, R.A., Paule, R.C., Schaffer, R., et al., Standard Reference Materials: A Reference Method for the Determination of Potassium in Serum, NBS Spec. Publ. 260-63 (May 1979). PB297207**

- Velapoldi, R.A., and Mielenz, K.D., Standard Reference Materials: A Fluorescence SRM Quinine Sulfate Dihydrate (SRM 936), NBS Spec. Publ. 260-64 (January 1980). PB80132046**
- Marinenko, R.B., Heinrich, K.F.J., and Ruegg, F.C., Standard Reference Materials:
 Micro-Homogeneity Studies of NBS SRM, NBS Research Materials, and Other Related Samples, NBS Spec. Publ. 260-65 (September 1979). PB300461**
- Venable, W.H., Jr., and Eckerle, K.L., Standard Reference Materials: Didymium Glass Filters for Calibrating the Wavelength Scale of Spectrophotometers (SRMs 2009, 2010, 2013, 2014). NBS Spec. Publ. 260-66 (October 1979). PB80104961**
- Velapoldi, R.A., Paule, R.C., Schaffer, R., et al., Standard Reference Materials: A Reference Method for the Determination of Chloride in Serum, NBS Spec. Publ. 260-67 (November 1979). PB80110117**
- Mavrodineanu, R., and Baldwin, J.R., Standard Reference Materials: Metal-On-Quartz Filters as a SRM for Spectrophotometry SRM 2031, NBS Spec. Publ. 260-68 (April 1980). PB80197486**
- Velapoldi, R.A., Paule, R.C., Schaffer, R., et al., Standard Reference Materials: A Reference Method for the Determination of Lithium in Serum, NBS Spec. Publ. 260-69 (July 1980). PB80209117**
- Marinenko, R.B., Biancaniello, F., Boyer, P.A., et al., Standard Reference Materials: Preparation and Characterization of an Iron-Chromium-Nickel Alloy for Microanalysis: SRM 479a, NBS Spec. Publ. 260-70 (May 1981). SN003-003-02328-1*
- Seward, R.W., and Mavrodineanu, R., Standard Reference Materials: Summary of the Clinical Laboratory Standards Issued by the National Bureau of Standards, NBS Spec. Publ. 260-71 (November 1981). PB82135161**
- Reeder, D.J., Coxon, B., Enagonio, D., et al., Standard Reference Materials: SRM 900, Antiepilepsy Drug Level Assay Standard, NBS Spec. Publ. 260-72 (June 1981). PB81220758

- Interrante, C.G., and Hicho, G.E., Standard Reference Materials: A Standard Reference Material Containing Nominally Fifteen Percent Austenite (SRM 486), NBS Spec. Publ. 260-73 (January 1982). PB82215559**
- Marinenko, R.B., Standard Reference Materials: Preparation and Characterization of K-411 and K-412 Mineral Glasses for Microanalysis: SRM 470, NBS Spec. Publ. 260-74 (April 1982). PB82221300**
- Weidner, V.R., and Hsia, J.J., Standard Reference Materials: Preparation and Calibration of First Surface Aluminum Mirror Specular Reflectance Standards (SRM 2003a), NBS Spec. Publ. 260-75 (May 1982). PB82221367**
- Hicho, G.E., and Eaton, E.E., Standard Reference Materials: A Standard Reference Material Containing Nominally Five Percent Austenite (SRM 485a), NBS Spec. Publ. 260-76 (August 1982). PB83115568**
- Furukawa, G.T., Riddle, J.L., Bigge, W.G., et al., Standard Reference Materials: Application of Some Metal SRMs as Thermometric Fixed Points, NBS Spec. Publ. 260-77 (August 1982). PB83117325**
- Hicho, G.E., and Eaton, E.E., Standard Reference Materials: Standard Reference Material Containing Nominally Thirty Percent Austenite (SRM 487), NBS Spec. Publ. 260-78 (September 1982). PB83115576**
- Richmond, J.C., Hsia, J.J., Weidner, V.R., et al., Standard Reference Materials: Second Surface Mirror Standards of Specular Spectral Reflectance (SRMs 2023, 2024, 2025), NBS Spec. Publ. 260-79 (October 1982). PB84203447**
- Schaffer, R., Mandel, J., Sun, T., et al., Standard Reference Materials: Evaluation by an ID/MS Method of the AACC Reference Method for Serum Glucose, NBS Spec. Publ. 260-80 (October 1982). PB84216894**
- Burke, R.W., and Mavrodineanu, R., Standard Reference Materials: Accuracy in Analytical Spectrophotometry, NBS Spec. Publ. 260-81 (April 1983). PB83214536**

- Weidner, V.R., Standard Reference Materials: White Opal Glass Diffuse Spectral Reflectance Standards for the Visible Spectrum (SRMs 2015 and 2016), NBS Spec. Publ. 260-82 (April 1983). PB83220723**
- Bowers, G.N., Jr., Alvarez, R., Cali, J.P., et al., Standard Reference Materials: The Measurement of the Catalytic (Activity) Concentration of Seven Enzymes in NBS Human Serum (SRM 909), NBS Spec. Publ. 260-83 (June 1983). PB83239509**
- Gills, T.E., Seward, R.W., Collins, R.J., et al., Standard Reference Materials: Sampling, Materials Handling, Processing, and Packaging of NBS Sulfur in Coal SRMs 2682, 2683, 2684, and 2685, NBS Spec. Publ. 260-84 (August 1983). PB84109552**
- Swyt, D.A., Standard Reference Materials: A Look at Techniques for the Dimensional Calibration of Standard Microscopic Particles, NBS Spec. Publ. 260-85 (September 1983). PB84112648**
- Hicho, G.E., and Eaton, E.E., Standard Reference Materials: A SRM Containing Two and One-Half Percent Austenite, SRM 488, NBS Spec. Publ. 260-86 (December 1983). PB84143296**
- Mangum, B.W., Standard Reference Materials: SRM 1969: Rubidium Triple-Point - A Temperature Reference Standard Near 39.30° C, NBS Spec. Publ. 260-87 (December 1983). PB84149996**
- Gladney, E.S., Burns, C.E., Perrin, D.R., et al., Standard Reference Materials: 1982 Compilation of Elemental Concentration Data for NBS Biological, Geological, and Environmental Standard Reference Materials, NBS Spec. Publ. 260-88 (March 1984). PB84218338**
- Hust, J.G., Standard Reference Materials: A Fine-Grained, Isotropic Graphite for Use as NBS
 Thermophysical Property RMs from 5 to 2500
 K, NBS Spec. Publ. 260-89 (September 1984).
 PB85112886**
- Hust, J.G., and Lankford, A.B., Standard Reference Materials: Update of Thermal Conductivity and Electrical Resistivity of Electrolytic Iron, Tungsten, and Stainless Steel, NBS Spec. Publ. 260-90 (September 1984). PB85115814**

- Goodrich, L.F., Vecchia, D.F., Pittman, E.S., et al., Standard Reference Materials: Critical Current Measurements on an NbTi Superconducting Wire SRM, NBS Spec. Publ. 260-91 (September 1984). PB85118594**
- Carpenter, B.S., Standard Reference Materials:
 Calibrated Glass Standards for Fission Track Use
 (Supplement to NBS Spec. Publ. 260-49), NBS
 Spec. Publ. 260-92 (September 1984).
 PB85113025**
- Ehrstein, J.R., Standard Reference Materials:
 Preparation and Certification of SRM for
 Calibration of Spreading Resistance Probes, NBS
 Spec. Publ. 260-93 (January 1985).
 PB85177921**
- Gills, T.E., Koch, W.F., Stolz, J.W., et al., Standard Reference Materials: Methods and Procedures Used at the National Bureau of Standards to Certify Sulfur in Coal SRMs for Sulfur Content, Calorific Value, Ash Content, NBS Spec. Publ. 260-94 (December 1984). PB85165900**
- Mulholland, G.W., Hartman, A.W., Hembree, G.G., et al., Standard Reference Materials: Development of a 1mm Diameter Particle Size Standard, SRM 1690, NBS Spec. Publ. 260-95 (May 1985). PB95-232518/AS**
- Carpenter, B.S., Gramlich, J.W., Greenberg, R.R., et al., Standard Reference Materials: Uranium-235 Isotopic Abundance Standard Reference Materials for Gamma Spectrometry Measurements, NRS Spec. Publ. 260-96 (September 1986). PB87108544**
- Mavrodineanu, R., and Gills, T.E., Standard Reference Materials: Summary of the Coal, Ore, Mineral, Rock, and Refactory Standards Issued by the National Bureau of Standards, NBS Spec. Publ. 260-97 (September 1985). PB86110830**
- Hust, J.G., Standard Reference Materials: Glass Fiberboard SRM for Thermal Resistance, NBS Spec. Publ. 260-98 (August 1985). SN003-003-02674-3*
- Callanan, J.E., Sullivan, S.A., and Vecchia, D.F., Standard Reference Materials: Feasibility Study for the Development of Standards Using Differential Scanning Calorimetry, NBS Spec. Publ. 260-99 (August 1985). PB86106747**

- Taylor, J.K., Trahey, N.M., ed., Standard Reference Materials: Handbook for SRM Users, NBS Spec. Publ. 260-100 (February 1993). PB93183796**
- Mangum, B.W., Standard Reference Materials: SRM 1970, Succinonitrile Triple-Point Standard: A Temperature Reference Standard Near 58.08° C, NBS Spec. Publ. 260-101 (March 1986). PB86197100**
- Weidner, V.R., Mavrodineanu, R., Mielenz, K.D., et al., Standard Reference Materials: Holmium Oxide Solution Wavelength Standard from 240 to 640 nm SRM 2034, NBS Spec. Publ. 260-102 (July 1986). PB86245727**
- Hust, J.G., Standard Reference Materials: Glass Fiberblanket SRM for Thermal Resistance, NBS Spec. Publ. 260-103 (September 1985). PB86109949**
- Mavrodineanu, R., and Alvarez, R., Standard Reference Materials: Summary of the Biological and Botanical Standards Issued by the National Bureau of Standards, NBS Spec. Publ. 260-104 (November 1985). PB86155561**
- Mavrodineanu, R., and Rasberry, S.D., Standard Reference Materials: Summary of the Environmental Research, Analysis, and Control Standards Issued by the National Bureau of Standards, NBS Spec. Publ. 260-105 (March 1986). PB86204005**
- Koch, W.F., ed., Standard Reference Materials: Methods and Procedures Used at the National Bureau of Standards to Prepare, Analyze, and Certify SRM 2694, Simulated Rainwater, and Recommendations for Use, NBS Spec. Publ. 260-106 (July 1986). PB86247483**
- Hartman, A.W., and McKenzie, R.L., Standard
 Reference Materials: SRM 1965, Microsphere
 Slide (10 μm Polystyrene Spheres), NIST Spec.
 Publ. 260-107 (November 1988).
 PB89153704**
- Mavrodineanu, R., and Gills, T.E., Standard Reference Materials: Summary of Gas Cylinder and Permeation Tube Standard Reference Materials Issued by the National Bureau of Standards, NBS Spec. Publ. 260-108 (May 1987). PB87209953**

- Candela, G.A., Chandler-Horowitz, D., Novotny, D.B., et al., Standard Reference Materials: Preparation and Certification of an Ellipsometrically Derived Thickness and Refractive Index Standard of a Silicon Dioxide Film (SRM 2530), NIST Spec. Publ. 260-109 (October 1988). PB89133573**
- Kirby, R.K., and Kanare, H.M., Standard Reference Materials: Portland Cement Chemical Composition Standards (Blending, Packaging, and Testing), NBS Spec. Publ. 260-110 (February 1988). PB88193347**
- Gladney, E.S., O'Malley, B.T., Roelandts, I., et al., Standard Reference Materials: Compilation of Elemental Concentration Data for NBS Clinical, Biological, Geological, and Environmental Standard Reference Materials, NBS Spec. Publ. 260-111 (November 1987). PB88156708**
- Marinenko, R.B., Blackburn, D.H., and Bodkin,
 J.B., Standard Reference Materials: Glasses for
 Microanalysis: SRMs 1871-1875, NIST Spec.
 Publ. 260-112 (February 1990). PB90215807**
- Mangum, B.W., and Wise, J.A., Standard Reference Materials: Description and Use of a Precision Thermometer for the Clinical Laboratory, SRM 934, NIST Spec. Publ. 260-113 (June 1990). PB90257643**
- Vezzetti, C.F., Varner, R.N., and Potzick, J.E., Standard Reference Materials: Bright-Chromium Linewidth Standard, SRM 476, for Calibration of Optical Microscope Linewidth Measuring Systems, NIST Spec. Publ. 260-114 (January 1991). PB91167163**
- Williamson, M.P., Willman, N.E., and Grubb, D.S., Standard Reference Materials: Calibration of NIST SRM 3201 for 0.5 in. (12.65 mm) Serial Serpentine Magnetic Tape Cartridge, NIST Spec. Publ. 260-115 (February 1991). PB91187542**
- Mavrodineanu, R., Burke, R.W., Baldwin, J.R., et al., Standard Reference Materials: Glass Filters as a Standard Reference Material for Spectrophotometry-Selection, Preparation, Certification and Use of SRM 930 and SRM 1930, NIST Spec. Publ. 260-116 (March 1994). PB94-188844/AS**

- Vezzetti, C.F., Varner, R.N., and Potzick, J.E., Standard Reference Materials: Antireflecting-Chromium Linewidth Standard, SRM 475, for Calibration of Optical Microscope Linewidth Measuring Systems, NIST Spec. Publ. 260-117 (January 1992). PB92-149798**
- Williamson, M.P., Standard Reference Materials: Calibration of NIST Standard Reference Material 3202 for 18-Track, Parallel, and 36-Track, Parallel Serpentine, 12.65 mm (0.5 in), 1491 cpmm (37871 cpi), Magnetic Tape Cartridge, NIST Spec. Publ. 260-118 (July 1992). PB92-226281**
- Vezzetti, C.F., Varner, R.N., and Potzick, Standard Reference Materials: Antireflecting-Chromium Linewidth Standard, SRM 473, for Calibration of Optical Microscope Linewidth Measuring System, NIST Spec. Publ. 260-119 (September 1992)
- Caskey, G.W., Philips, S.D., Borchardt., et al., Standard Reference Materials: A Users' Guide to NIST SRM 2084: CMM Probe Performance Standard, NIST Spec. Publ. 260-120 (1994)
- Rennex, B.G., Standard Reference Materials:
 Certification of a Standard Reference Material
 for the Determination of Interstitial Oxygen
 Concentration in Semiconductor Silicon by
 Infrared Spectrophotometry, NIST Spec. Publ.
 260-121 (1994) PB95-125076/AS
- Gupta, D., Wang, L., Hanssen, L.M., Hsai, J.J., and Datla, R.U., Polystyrene Films for Calibrating the Wavelength Scale of Infrared Spectrophotometer (SRM 1921). NIST Spec. Publ. 260-122 (1995) PB95-226866/AS
- Development of Technology and the Manufacture of Spectrometric SRMs for Naval Brasses (MC62 M63). NIST Spec. Publ. 260-123 (IN PREP).
- Strouse, G.F., SRM 1744: Aluminum Freezing Point Standard. NIST Spec. Publ. 260-124 (1995) SN003-003-03342-1
- Schiller, S.B, Standard Reference Materials: Statistical Aspects of the Certification of Chemical Batch SRMs. NIST Spec. Publ. 260-125 (1996) PB96-210877/AS

- Guenther, F.R., Dorko, W.D., Miller, W.R., et al., Standard Reference Materials: The NIST Traceable Reference Material Program for Gas Standards, NIST Spec. Publ. 260-126 (1996) PB96-210786/AS
- Strouse, G.F., and Ahmet, A.T., Standard Reference Material 1747: Tin Freezing-Point Cell and Standard Reference Material 1748: Zinc Freezing-Point Cell. NIST Spec. Publ. 260-127 (IN PREP).
- Zhang, Z.M., Gentile, T.R., Migdall, A.L., and Datla, R.U., Transmission Filters with Measured Optical Density at 1064 nm Wavelength--SRM 2036. SRM Spec. Publ. 260-128 (IN PREP).
- Potzick, J.E., Antireflecting-Chromium Linewidth Standard, Standard Reference Material 473, for Calibration of Optical Microscope Linewidth Measuring Systems, NIST Spec. Publ. 260-129 (1997) PB97-151922/AS.
- Zarr, R.R., Standard Reference Materials: Glass Fiberboard, Standard Reference Material 1450c, for Thermal Resistance from 280K to 340K, NIST Spec. Publ. 260-130 (IN PREP).

- *Send order with remittance to: Superintendent of Documents, U.S. Government Printing Office, Washington, DC 20402-9325. Remittance from foreign countries should include an additional one fourth of the purchase price for postage.
- **May be ordered from: National Technical Information Services (NTIS), Springfield, VA 22161.
 - For information phone (703-487-4650) To Place an Order with PB# phone (800-553-6847)

Table of Contents

	F F	Page
Al	bstract	. 1
Int	troduction	. 2
1.	Certification of Improved Resistivity SRMs 1.1 General Comments	. 3 . 5 . 6 . 7
2.	Certification Procedure	. 9 11
3.	Control Procedures for the Certification Process	14 15
4.	Equations Used for Calculating Sheet Resistance and Resistivity Values 4.1 Rewriting the Equation to Relate to Evaluation of Uncertainty	
5.	Sources of Measurement Uncertainty — Detailed Discussion	19
	Average Value	21 21 22 25
	Measurements	30 33
	Factor	34

Table of Contents (cont'd.)

	P	age
6.	Unanticipated Effects	36
7.	Compilation of Uncertainty Components	39 40 42
8.	Conclusion	44
9.	Acknowledgments	44
Re	ferences	44
Ap Ap Ap Ap	pendix 1. Summary of Important SRM Wafer Material and Measurement Condition Parameters	47 66 70 74 78
	List of Figures	
	P	age
1.	Scaled drawing of a 100 mm diameter wafer showing locations of thickness measurements and the locations of the four-point probe measurements in the 2X magnification at the bottom	10
2.	Schematic of probe wiring for dual-configuration measurements	12
3.	Conceptual drawing of wafer with two textured surfaces during thickness	32

List of Tables

	Page
1.	Manufacturers' Specifications for the Current Supplies and DVM Used for Certification
2.	Standard Resistor Values, and Typical Measurement Voltages for Each of the SRM Levels
3.	Temperature Coefficients of Resistivity of Silicon for the Nominal Values of the SRMs
4.	Variance in Resistivity Value Due to Temperature Error 30
5.	Specifications for Haidenhain Certo 60 Thickness Measurement Instrument
6.	Components Identified in Statistical Analyses of Certification and Control Experiment Data
7.	Values of the Components Identified in Statistical Analyses for the Various SRMs in Appendices 2 through 6, $m\Omega$ cm
8.	Type A Standard Uncertainty Values, u_i , Taken from Appendices 2 through 6, m Ω cm 40
9.	Type A Variance Values, u_i^2 , Obtained by Squaring the Entries in Table 8 41
10.	Type B Variance Values, u_j^2 , Calculated from Summation of Terms in Sections 5.2.1 through 5.2.4
11.	Combined Variance Values, u_c^2 , from Addition of Terms from Table 9 and 10 42
12.	Combined Standard Uncertainty Values, u_c
13.	Expanded Uncertainty Values, U
14.	Bias Corrections Applied to Measured Values

Standard Reference Materials: The Certification of 100 mm Diameter Silicon Resistivity SRMs 2541 through 2547 Using Dual-Configuration Four-Point Probe Measurements

by

James R. Ehrstein Semiconductor Electronics Division, EEEL

and

M. Carroll Croarkin Statistical Engineering Division, ITL

ABSTRACT

This report documents the selection of material, the certification procedure and its control, and the analysis of measurement uncertainty for a family of new and improved Standard Reference Materials for sheet resistance and resistivity of silicon wafers, SRMs 2541 through 2547, covering the resistivity range $0.01~\Omega$ cm through 200 Ω cm. These SRMs, made from 100 mm silicon, replace previous SRM sets 1521 through 1523, which used 50.8 mm (2 in) diameter silicon at the same nominal resistivity levels.

The certification of the new SRMs uses a dual-configuration four-point probe procedure rather than the single-configuration procedure of ASTM F84, as used for previous SRMs. The new SRMs offer better handling compatibility with current user instrumentation, better uniformity of wafer thickness and of resistivity, more extensive spatial characterization of the near-center wafer resistivity, and reduced measurement uncertainty compared to the SRMs they replace.

The general procedures for the certification measurements, the control of the certification process, and the analysis of the results are based on experience gained from numerous preliminary experiments that allowed evaluation of the importance and relative magnitude of many possible measurement effects. The validity and effectiveness of the resulting certification and control procedures were tested during the analysis of results from the first of the SRMs to be certified, that at 200 Ω -cm. The body of this report details the background and principles of the certification process and the approach to analyzing the experimental data needed to calculate the uncertainty of the certified values. This report details the evaluation of underlying components of uncertainty that apply to all SRM levels. Additional components derived from statistical analyses of the actual certification data are done separately for each SRM level and are reported in the appendices for all SRMs except SRM 2543

and SRM 2544. The analyses of certification for these latter two SRMs have not been completed at this time and will be documented in supplements to this report when available.

Key Words: four-point probe; resistivity; semiconductor; silicon; SRM; standards

INTRODUCTION

This Special Publication summarizes the certification procedure for a new generation of silicon resistivity Standard Reference Materials (SRMs) 2541 through 2547. It includes in individual appendices the analysis of the associated uncertainty levels calculated from the certification data for each of the resistivity levels except for SRMs 2543 and 2544, the statistical analyses for which will be given in a supplementary report.

Previous Resistivity SRMs

For a number of years, the Semiconductor Electronics Division of NIST has issued three sets of silicon resistivity SRMs. These sets, designated 1521, 1522, and 1523, contain two, three, and two wafers, respectively, of 50.8 mm (2 in) diameter silicon with the resistivity values in each set having been chosen to serve a particular application need in the silicon semiconductor industry. The generic purpose of each of these sets was to allow a user to verify the performance of a four-point probe test instrument, or to calibrate the output of a noncontact eddy current conductance measuring instrument. The range of resistivity values of these SRMs is from about 0.01 Ω cm to about 200 Ω cm. A total of more than 1300 of these sets has been certified and sold worldwide. Each wafer in each set was certified for resistivity using a four-point probe following the measurement procedure of ASTM Method F84 [1]. This procedure is also referred to as a "single-configuration" four-probe procedure in the remainder of this report.

Improved Resistivity SRMs

After several years of exploratory work, the Semiconductor Electronics Division is issuing improved SRMs at the same resistivity levels as in the previous sets, but having four salient upgraded features:

1. The new SRMs are wafers of 100 mm diameter silicon which enables better compatibility with present generation user instrumentation.

^{*}In the remainder of this report, the term "four-point probe" is used when referring to the probe itself. The term "four-probe" is used when referring to the measurement process.

- 2. They are fabricated from silicon with improved uniformity of resistivity and thickness. This will reduce ambiguities of interpretation related to measurement sampling volume, and it will improve transferability of the certified value to the end user.
- 3. They are certified with a modification of the original certification procedure. This modified procedure is referred to as "dual-configuration" or "configuration-switched" four-probe measurements and is implemented on most commercial four-point probe instruments that are automated for thin-film sheet resistance mapping. Tests at NIST have shown there is also a significant reduction of uncertainty when using this procedure for measuring bulk silicon wafers.
- 4. Measurements and analysis are provided at the wafer center, as was done with the original SRMs, and also around two small circles with sizes related to the requirements of commercial resistivity-measuring instrumentation. These additional data serve to characterize the small nonuniformities in resistivity that are present even in these wafers.

While these improved SRMs are issued singly, rather than in sets, it is strongly recommended that for all purposes of calibration or testing of instrument linearity, SRMs at two or more resistivity levels be used, with the values being chosen according to the user's application needs.

A major goal of these SRMs has been to meet or exceed the requirements set forth at the SEMATECH Workshop on Silicon Materials for Mega-IC Applications [2]: "That layer resistivity measurements be improved to an accuracy of 1 % and repeatability of 0.5 % and that NIST provide the SRMs required for such measurements."

1. CERTIFICATION OF IMPROVED RESISTIVITY SRMs

1.1 General Comments

While it may seem trivial to generate silicon resistivity standards that far exceed the SEMATECH Mega-IC Workshop requirements for precision and accuracy, this is not the case. Silicon is a semiconductor, nonuniform in resistivity in both lateral and vertical directions, unpassivated for use as an SRM, that can be measured with a four-point probe and, therefore, subject to possible surface effects due to storage and handling environments that can modify the near-surface resistivity. A lapped surface is used on SRM wafers to increase surface recombination velocity, to improve the quality of contact with the spring-loaded probe tips, and to improve the long-term stability of measured resistivity by reducing the susceptibility to changing surface conditions. This, in turn, introduces compromises in terms of near-surface damage, and of the definition and measurement of wafer thickness. Four-point probes are used for certification measurements, and the probes are subject to wear and to changes in contact quality and performance that may be either gradual or rather sudden. Despite a number of efforts, no simple characteristic of a probe pin has been identified that is

a clear indicator of how that pin will contribute to the quantitative performance of a given four-point probe. It has been found that measurement precision with a single probe head, as well as measurement variability among probe heads, are functions (among other things) of resistivity, conductivity type, specimen surface preparation, environmental conditions, and present condition of the probe pins themselves.

It is possible, using a technique such as the van der Pauw procedure [3] with contacts bonded to the perimeter of a polished wafer and with measurements done in an ambient capable of controlling wafer surface charge, to eliminate many of the concerns related to measurements with mechanical probe contacts. It might be possible in this way to eliminate or reduce noticeably a number of sources of measurement variability. Such measurements would then have a lower uncertainty than those made by four-point probe on a lapped wafer and might well provide the best estimate of overall volume average resistivity for an entire silicon wafer. However, this would probably not be particularly useful for calibrating or verifying the performance of instruments used in production environments if the measurements required a special ambient for measurement or if the full-wafer average resistivity did not bear a clear relationship to the localized (small area) value measured by the production test instruments.

An important distinction needs to be made. The principal objective of these SRMs is not to provide the best value of the volume resistivity of the silicon wafer itself, but to use the SRM wafer to help define and transfer a functional resistivity/sheet resistance measurement scale to users of common instrumentation in various parts of the semiconductor industry. Currently, most such equipment is based on four-probe dc resistance or on eddy current measurements and has spatial sampling volumes on the centimeter scale. There is no known analytic expression for the exact volume weighting of measurements by a four-point probe or by an eddy current tester with a ferrite core. As a result, it is not possible to guarantee perfect equivalence between four-probe and eddy current instruments for specimens with various and arbitrary patterns of resistivity nonuniformity. Nevertheless, resistivity SRMs based on lapped silicon wafers with certification measurements by four-point probe, particularly when done with a well-controlled measurement system used in the dual-configuration mode, and with measurements in well-specified locations on the SRM wafer, offer the user community several significant benefits. These are: stable SRM artifacts, measurement sampling volume generally comparable to that of the user's instrumentation, and certified measurement precision and resolution that more than meets the requirements of the semiconductor industry. Thus, in developing these SRMs, the interest is not so much in what the true bulk resistivity of each silicon wafer is, but rather in how the measurement values on these wafers behave as a function of measurement conditions, and how the SRMs transfer between NIST and the user community.

There are two principal reasons for preferring the dual-configuration implementation of four-probe measurements for the SRM certification. First, the probe-to-probe differences are reduced noticeably compared to those that exist when using the single-configuration (ASTM) procedure; such differences are generally only several tenths of a percent, but make it difficult to reach or exceed the accuracy goals in the SEMATECH Mega-IC Workshop report.

Second, the scatter, or random error, is reduced in a set of measurements taken with any given probe. Both improvements are interpreted as being due to the ability of dual-configuration measurements to correct more exactly for the true electrical probe separations than can be done with the auxiliary optical and mechanical separation measurements required by the ASTM Method, in combination with single-configuration electrical data.

For some time, it was common among users of the technique to speak of configuration-switched rather than dual-configuration four-probe measurements. The term dual configuration is used in this report when the term is written out in order to reinforce the operational difference from the ASTM, or single-configuration, procedure.

The following sections discuss the details of the procedures used for certification, and its control, as well as the manner of analyzing, and reporting the results. They also give a brief description of the components of measurement uncertainty in relation to the equation used to calculate the reported results from the raw data. Section 5 discusses the evaluation of uncertainty in more detail, but organizes the discussion according to whether the various contributions are evaluated by ISO Type A or Type B evaluation procedures (see 1.3).

1.2 Resistivity Standards vs. Sheet Resistance Standards

This SRM is called a "resistivity" standard, and much of this report and the SRM certificate focus on describing it and analyzing the measurements in terms of a resistivity value. This is done primarily as a concession to customary terminology and conceptualization in the semiconductor industry wherein "sheet resistance" is a property associated almost exclusively with a thin film of conducting material rather than with a substrate wafer. However, these SRM wafers do have sheet resistance values associated with them (resistivity divided by thickness), and moreover, the functional need of most user instrumentation is actually for calibration or verification of a (sheet) resistance scale, and not of a resistivity scale.

This distinction is not simply one of semantics. There is an actual benefit to the user from treating the SRMs as sheet resistance reference artifacts. To obtain resistivity values for a silicon wafer, it is necessary to know the wafer's thickness. But when silicon wafers are lapped in order to improve their stability as electrical reference standards, the surface texture compromises the possibility of a wafer having a single, unique wafer thickness. The determination of the wafer's resistivity value is therefore poorer than that of its sheet resistance value because of the added uncertainty due to thickness. Thus, each of the SRM wafers has a somewhat larger relative uncertainty of resistivity than it does for sheet resistance.

Further, if the user employs these SRMs to establish a scale for resistivity but uses an independent measurement of thickness such as from a capacitive- or sonic-gauge, then the user must add yet another component of uncertainty to the transfer process. The reason is that these other instrument types are different in operating principle from that of the

contacting electronic-micrometer which is used at NIST for the determination of SRM wafer thicknesses. Therefore, they are not likely to give the same functional value of wafer thickness that is reported on the SRM certificate, and an additional measurement error is incurred in establishing a resistivity scale. However, sheet resistance values do not depend on measured thickness value, and transfer of SRM (sheet) resistance values are unaffected by this consideration. It is therefore recommended: 1) that these SRMs be used as **sheet resistance standards** whenever possible and 2) that the thickness value given on the certificate be used whenever a resistivity value is needed.

[Note: Thickness values for lapped surface wafers typical of those being used for these SRMs have been found to be about 0.5 % smaller when measured with a capacitance gauge than when measured with an electronic-micrometer. These capacitance-gauge thickness values are probably closer to the actual thickness of the electrically conducting portion of the wafer (beneath the lapped texture) than are those from the electronic-micrometer. However, for the purposes of SRM certification, it is easier to establish traceability of thickness scale to dimensional standards when using an electronic-micrometer.]

1.3 Traditional Description of Uncertainty and the ISO Formulation

Measurement uncertainty for these SRMs is reported in conformance with guidelines formulated by the International Standards Organization, ISO [4, 5]. Sources of uncertainty are classified as Type A or Type B according to whether their values are estimated from repeated measurements (Type A), or are inferred in another manner (Type B). A variance is calculated, or estimated, for each contribution to the uncertainty of the measured value; a sum of variances is then done separately for Type A and Type B evaluations. The square root of the sum of the Type A and Type B variances is calculated and is called the combined standard uncertainty, u_c . A quantity called the expanded uncertainty, U_c , is calculated by multiplying the standard uncertainty by a coverage factor, k. This factor can often be taken from the Student t tables to give a stated coverage, say 95 %, if the degrees of freedom can be calculated. The effective degrees of freedom in the analyses of each of the SRM levels are sufficiently large, typically 60 or more, that a factor of k = 2 gives a coverage of 95 %.

Where sources of uncertainty for this SRM are estimated from other than repeated measurements, it is generally assumed that the affected measurements come from a rectangular distribution, the limits of which are the values that would have been assumed as the maximum systematic error for that quantity. For a rectangular distribution, the variance is the half-width divided by $\sqrt{3}$. There is not always a one-to-one correspondence between the categorization of traditional sources of measurement error as being random or systematic and the uncertainty components determined by Type A or Type B evaluation procedures.

1.4 Acquisition and Characteristics of Silicon Wafers for the SRMs

Wafers at all SRM resistivity levels were bought, having been already cut, etched, and lapped by the supplier. The supplier for each of the resistivity levels is identified on the SRM certificate. All wafers are nominally 625 μ m thick. The perimeters of all wafers were contoured to reduce breakage; a single primary orientation flat was ground onto all crystals prior to slicing. The supplier of the wafers for the three lowest resistivity levels, 0.01 Ω cm, 0.1 Ω cm, and 1 Ω cm, used a laser marking technique to engrave a unique wafer identification into each wafer just above this flat; the suppliers of the four highest levels did not offer such a marking process.

Wafers at the three lowest resistivity levels are from (100) boron-doped Czochralski-process (Cz) silicon crystals, while wafers at the four highest levels are from (111) crystals phosphorus-doped by the neutron-transmutation doping (NTD) process. These combinations have been found to be appropriate for meeting the goal of high uniformity of resistivity across a resulting wafer.

The suppliers** selected (Recticon Inc. for the three lowest resistivity levels, Wacker Siltronic for the middle level, and Topsil Semiconductor Materials A/S for the three highest levels) specialize in the types of growth processes listed. Preliminary batches of wafers from each supplier were evaluated for thickness and resistivity uniformity. These evaluations indicated a high degree of likelihood of total thickness variation being less than 1 µm over the wafer surface and of resistivity uniformity being 1 % or better within the central 50 mm diameter of the wafers. These levels of uniformity are not guaranteed, however.

1.5 Measurement Concerns and Control of the Certification Procedure

Extensive preliminary testing was done to reach a reasonable optimization of the wafer preparation and test conditions, to minimize or eliminate effects that would degrade the certification uncertainty, and to estimate the relative importance of the various known remaining effects. These tests then led to the design of several experimental control procedures to monitor and evaluate possible changes in probes, wafers, or instrumentation during the certification.

The following sources of experimental variability and possible error were identified and are listed along with the procedure that was developed to minimize their effect and to estimate their value.

^{**}Certain commercial equipment, instruments, or materials are identified in this report to specify adequately the experimental procedure. Such identification does not imply recommendation or endorsement by the National Institute of Standards and Technology, nor does it imply that the materials or equipment identified are necessarily the best available for the purpose.

- 1. Short-term imprecision (repeatability of measurements taken within a period of minutes in a small, uniform region of material) is believed to be controlled by probe contact fluctuations and electronics noise; it is minimized by using the dual-configuration procedure and choosing a probe with low noise. Short-term imprecision is evaluated from data at the centers of the certified wafers, as well as from all wafers used in the control procedures.
- 2. Longer-term imprecision (the ability to reproduce an average value at a fixed point on a wafer over a period of days or weeks) is related to changes in the measurement environment, e.g., power-line conditions, electromagnetic interference, or humidity. No measurements are taken at a relative humidity above 50 %, and residual long-term imprecision can be evaluated from the control experiments.
- 3. Probe-to-probe differences in measured value have been seen to exist. Although small (0.1 %, or less), it is necessary to identify and select for certification a probe with low bias. This is done through the design of one of the control experiments. Residual offset for the selected probe is estimated through analysis of this control experiment data, and a correction applied to the measured results if the offset is statistically significant.
- 4. Possible drift of the measurement process with time, whether due to changes in the probe used for certification, the wafers being tested, or to strong changes in the measurement environment. Drift can be estimated from the design of one of the control experiments.
- 5. Possible dependence of the measured resistivity value on the current value is controlled by a very tight procedure for selecting the current level for each SRM wafer.
- 6. Wafer nonuniformity effects on the certified values of resistivity are minimized by using very high uniformity wafers and by using a tightly controlled procedure for selecting the measurement locations.
- 7. Error related to the temperature dependence of resistivity value is controlled by measuring the temperature of the wafer stage and applying a correction for the difference between ambient and a reference temperature of 23 °C for each line of measured data. An estimate is made of the uncertainty of the temperature correction, and this estimate is part of the Type B standard uncertainty.
- 8. Possible error related to the accuracy of the measurement current supply and the digital voltmeter (DVM) are minimized by using standard resistors to measure the current value and by using the same scale of the DVM for measurement of both

wafer voltage and standard resistor voltage drops. Residual uncertainty related to the voltage measurements is estimated by Type B procedures.

9. Possible error related to the accuracy of the thickness measurement tool is minimized by instrument checks, several times a day, on NIST-traceable gauge blocks of thicknesses very close to those of the wafers. All wafers with a total indicated runout in excess of 1 μm over a nine-point thickness measurement pattern are rejected. Residual thickness measurement uncertainty is estimated by Type B procedures.

Accumulated probe damage in the wafers should not be detectable within the duration of the tests being performed. Previous tests on approximately a dozen wafers similar to these SRMs showed no effect out to 3000 probings for most wafers. However, a few wafers in those tests did show noticeable shifts (about 5 %) in average resistivity and greatly reduced measurement precision after about 1500 probings.

2. CERTIFICATION PROCEDURE

To minimize the effects of test instrument performance on measurement accuracy, a high degree of reliance is placed on ratioing techniques for both wafer thickness and electrical measurements, with the instruments being checked against precision calibration standards. Thus, the instrument used for wafer thickness measurement is regularly verified against gauge blocks having thicknesses very close to those of the SRM wafers, and measurements of the voltage drops across the silicon wafer and the standard resistor are read on the same scale of the same digital voltmeter. The standard resistors employed for monitoring the current serve as the primary reference point for all electrical measurement values.

2.1 Wafer Thickness Screening and Thickness Measurement

Preliminary screening with a capacitance-type thickness instrument of a small random selection of wafers from each of the actual SRM batches showed typical within-wafer thickness variation to be 0.2 μm , or less, for the central region where four-probe measurements are taken. This is noticeably better than the uniformity requirement of 1 % (which would be about 6.2 μm for the SRM wafers), as required by ASTM F84 for referee resistivity measurements.

For the certification procedure, thickness measurements of each wafer are taken on a three-row by three-column grid with a distance of 19 mm between the wafer center and the corners (Fig. 1). (The locations are approximate since the wafers are positioned manually.) This nine-site sampling gives a reasonable measure of the thickness and its variation in the area used for electrical measurements. Because the small contact area of the electronic-micrometer is more sensitive to local fluctuations due to variations in lapped surface texture, there is more

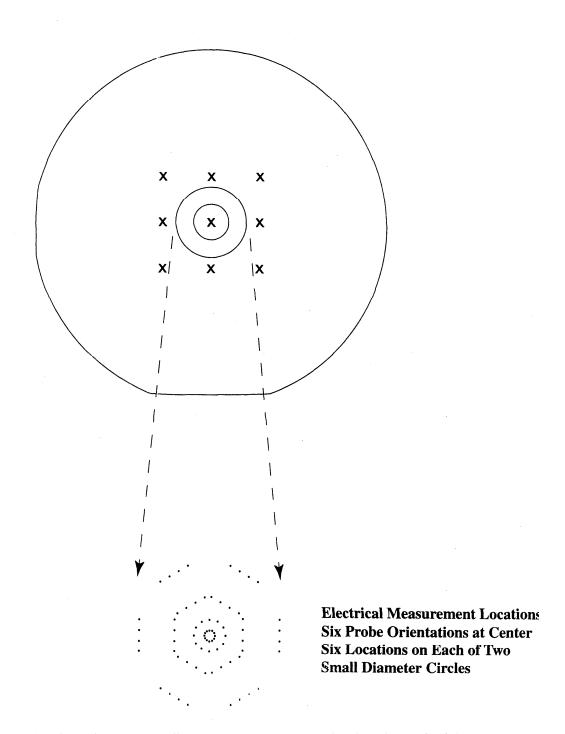


Figure 1. Scaled drawing of a 100 mm diameter wafer (top) showing locations of thickness measurements (x) and locations of the four-point probe measurements in the 2X magnification at the bottom.

variation in thickness values obtained by this instrument than those obtained by the capacitance gauge. Nevertheless, the range of thicknesses from this nine-point sampling plan is less than 0.5 µm for most wafers, and wafers are excluded from use as SRMs if there is an indicated variation of more than 1 µm among the nine sites. The average of all nine thickness values is used for conversion from sheet resistance to resistivity values on the SRM certificates. (ASTM Method F84 requires the use of only the thickness measured at the wafer center for this conversion.) The use of a nine-point average thickness reduces small errors due to local fluctuations in surface texture, is more representative of the area over which electrical measurements are taken, and improves the consistency among all wafers certified at a given SRM level. The standard deviation of these nine measurements is reported on each certificate for each wafer. Specifications for the electronic micrometer used for the thickness measurements can be found in Section 5.2.4.

2.2 Four-Probe Measurements of Sheet Resistance

Certification measurements are taken using a single four-point probe head, selected from five available (see 3.1). The specific probe used may differ from one resistivity level to another according to results of preliminary tests. All probe heads are constructed with in-line mounted tungsten-carbide probe pins, with a nominal separation of 1.59 mm between adjacent pins, with a spring-loaded force of about 1.5 N per pin and a nominal 40 µm (0.0016 in) tip radius. Eighteen sites are measured on each wafer, and wafers are allowed to equilibrate with the environment of the lab module for at least 24 h and with the temperature of the heat sink on the probe station for at least 1 min before being measured. Basic equipment requirements for all measurements follow ASTM Method F84; manufacturers' specifications for the equipment used can be found in Section 5.2.1. The measurement procedure for the first wiring configuration at each site follows ASTM F84, and that for the second configuration follows ASTM F1529 [6].

At each of the 18 sites for electrical measurement, the probe is connected first to the dc current source and DVM as in ASTM F84 (current through the outer probes and potential drop across the wafer measured with the inner probes). The current supply is set to give a specimen voltage drop between 9.95 mV and 10.05 mV for the forward current polarity at the first wafer-center measurement site. The current-supply controls remain set at this position for all remaining measurements on the wafer. The standard resistor for measurement of current value is chosen so that the voltage across the standard resistor is larger than that across the wafer. Applied current and specimen voltages are measured for both current polarities, and the average voltage-to-current ratio is calculated from these "forward" and "reverse" readings (to eliminate Seebeck voltages) [7]. Standard resistor and wafer voltages are recorded to a resolution of 0.1 µV. (More detail on the voltage measurements is given in Sec. 5.2.1.) While still in contact with the wafer, the probe head is connected to the current supply and DVM in the second wiring configuration, with the current passing between one outer probe pin and the nonadjacent interior pin, and the specimen voltages being measured with the two remaining pins. Again, forward and reverse direction current and wafer voltage values are measured, and an average voltage-to-current ratio is calculated. (See Fig. 2 for

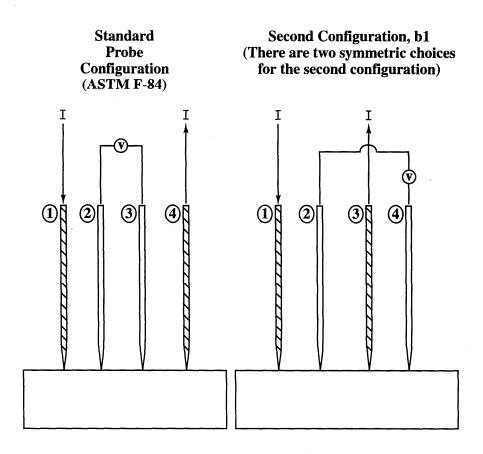


Figure 2. Schematic of probe wiring for dual-configuration measurements.

schematic of probe wiring.) For the wafers being certified, only one of the two nominally symmetric choices for this second wiring configuration is used, although both choices are used for measurement of the "control" wafers.

Using a theoretically derived relation between the voltage-to-current ratios from these two configurations, a scaling factor for lateral geometry effects is calculated [6]. This scaling factor, K_a , in eq (1) of Section 4, is multiplied by the voltage-to-current ratio from the first configuration to give a value for sheet resistance that has been corrected for wafer-edge boundary condition effects, and for variations of probe separation (at least to first order); an additional scaling factor is required if the wafer thickness is more than about 0.4 times the probe spacing (the largest ratio of wafer-thickness-to-probe-spacing for any of the SRM wafers is 0.4008). Multiplication of the sheet resistance values by wafer thickness produces values of wafer resistivity. The measurement results are corrected to 23 °C using the temperature of the heat sink at time of measurement and empirical temperature coefficients of resistivity [1].

A complete control and certification procedure is applied to a batch (approximately 125 wafers) at one of the seven resistivity levels; all data are analyzed for that batch and any necessary auxiliary measurements taken before proceeding to another resistivity level. This is to assure that any wear-induced drift, or other change that may be experienced by any of the four-point probes being used, is contained in and analyzed as part of the certification of a single batch. Two levels of control detailed in Sections 3.1 and 3.2 are used. In the first, and simpler part of certification control, a monitor-, or check-wafer, selected at random from the batch being certified, is measured at random times at least twice a day, during actual certification, to check for time-of-measurement effects due to factors other than changes in the probe. In the second part, a formal control experiment is conducted just prior to and just following the certification measurements. The entire cycle for initial control-wafer measurements, certification data for a batch of wafers, and final control-wafer measurements takes approximately 5 to 6 weeks.

2.3 Reporting of Data from 18 Measurements on Each Wafer

The 18 electrical measurement sites are distributed as follows: 1) six are located at the wafer center, with the wafer being rotated 30° between them, 2) six are spaced 60° apart around a circle of 5 mm (0.2 in) radius, and 3) six are spaced 60° apart around a circle of 10 mm (0.39 in) radius. (See Fig. 1.)

Average values of both sheet resistance and resistivity are reported for the center of the wafer, where wafer nonuniformity effects should be negligible. For measurements taken around the 5 mm and 10 mm radius circles, where additional variability due to material nonuniformity can be detected, individual site values are reported. To reduce clutter on the certificate, these individual values are given only for sheet resistance. A procedure for converting them to resistivity values follows eq (1), and is outlined on the certificate using values that are specific to each individual wafer. The data entries on the certificate are generally only

significant to several counts in the last digit. This digit is retained, however, to avoid additional error due to truncation.

The values reported for the two circles give the user a measure of the radial variation of resistivity for the wafer, although some azimuthal variation can also be detected on many wafers. This radial variation information is important for improving measurement transfer to instrument types having different integration volumes from that of the four-point probe used for certification. However, it is left to the user to determine how to weight the resistivities from the three regions of the wafer for the particular application of interest.

In cases where the user does not specifically need the resistance information from the 5 mm and 10 mm circles, it is strongly suggested that only the certified values from the wafer center be considered and that all user measurements be restricted to the wafer center.

3. CONTROL PROCEDURES FOR THE CERTIFICATION PROCESS

3.1 Control Procedure for Probe Effects

Immediately prior to certification of a batch of wafers and again at the end of certification, a "control" experiment is performed as follows. Each of five wafers, referred to as "controlwafers" and randomly selected from the batch to be certified, is measured for six orientations of the probe at the wafer center, using each of five probe heads in turn and the dual-configuration procedure. For the reason discussed in the next paragraph (criterion 4), both choices for the second wiring configuration are used for this test. This sequence is repeated until six rounds of measurements have been obtained on each wafer with each probe; the order of probes used and of the wafers measured is randomized for each round. Experience has shown that there is no reason to extend these measurements over a protracted period of time; this test is completed in about 7 days. The results are analyzed to give baseline values so the performance of the probe to be used for certification can be checked later if needed, to provide both short-term and longer-term estimates of measurement precision at that resistivity level, and to estimate the contribution to measurement uncertainty of the choice of measurement probe. For this latter purpose, the five available probes are assumed to represent a random sampling of all possible probes meeting reasonable operating requirements; they are not brought to like-new conditions prior to the tests. Probe heads are, however, prechecked for a number of operating characteristics, and individual pins replaced, if necessary.

After the results of the initial control experiment are analyzed, one of the probes is selected for certification measurements of all wafers in the batch. The following criteria are used when reviewing the initial control-wafer test data: 1) preference is given to probes with low average within-run standard deviations for six replicate runs on the five wafers; 2) preference is given to probes having high reproducibility of average value from the six rounds for each of the wafers; 3) preference is given to a probe that gives resistivity values in the middle of

the distribution for the six-round, five-probe, five-wafer data set which should ensure minimum bias to the ensuing certification data; and 4) preference is given to probes having good consistency of measured values between the two choices for wiring the second measurement configuration. Experience has shown that most probes do not give exactly the same measurement results on bulk substrate wafers for the two choices of second configuration. The small differences that are generally seen are believed due to the inability of dual-configuration measurements on bulk silicon to completely account for variations in probe spacing. Therefore, it is important to identify, and use for certification, probes that behave according to the theory for dual-configuration measurements where the theory does not admit to a distinction between the two choices for the second measurement configuration. There is no a priori formula or weighting factor used for these preference criteria; the goal is simply to identify and use the probe that has the lowest short-term and longer-term "noise" or imprecision and the least bias in measurement results.

This control experiment is repeated upon completion of the certification measurements for a wafer batch. This repetition is used to test for a change in response of the probe used for certification (which would indicate wear or contamination during certification). This should be distinguished from possible drift that might show up for most, or all, of the probes and which would more likely be due to changes in the measurement environment or to changes in the control-wafers themselves. Small changes in the response of only the certification probe would need to be accounted for by use of an additional contribution to the uncertainty statement for the SRM value. Larger changes in the response of the certification probe, if they occur, might require the probe to be rebuilt, and the entire sequence comprising initial control experiment, probe selection, batch certification, and final control experiment to be repeated. Changes in the response of all probes, if observed, would be analyzed for consistency or randomness of behavior and appropriate components estimated for the uncertainty statement.

Data from this multi-wafer control experiment also serve to estimate short-, intermediate-, and longer-term random variations in the certification process; see Appendix 2.

3.2 Control Procedure for Day-to-Day (Environmental) Effects

Acquisition of all the certification data on a batch of about 125 wafers takes approximately 10 to 12 days. Humidity is monitored, and no wafers are measured when relative humidity readings are in excess of 50 %. To monitor for possible effects due to changes in humidity, power-line fluctuations, or similar environmental problems, one wafer from the batch, referred to as a "monitor-wafer" or "check-wafer," is measured at random times approximately twice a day for the duration of certification. This results in 20 to 30 sets of measurements on the check-wafer (six wafer-center measurement sites each); this number is well below any level that has been found to cause significant change of value due to accumulated probing damage. These check-wafer data are analyzed for possible day-to-day (or time-of-day) variations in value, either random or systematic, that need to be incorporated into the uncertainty statement. The check-wafer data can also be used to give another

estimate of the short-term precision of the measurement process, and may serve to corroborate wear or contamination in the certifying probe.

3.3 Control Procedure for Other Longer Term Effects and Drift of Wafer or Probe

The sets of control-wafer measurements that are taken both before and after the certification data can be used to determine whether average resistivity has increased or decreased between the two series of measurements. If they have changed by a statistically significant amount, the multiplicity of probes involved can be used to determine whether the changes are likely due to wear of the certification probe (only that probe should show significant change) or whether the same changes are detected by most or all of the probes being used. The latter condition would indicate likely changes in the control wafers themselves or in the measurement equipment. Appropriate follow-up tests would then need to be made or suitable additional terms added to the uncertainty statement.

4. EQUATIONS USED FOR CALCULATING SHEET RESISTANCE AND RESISTIVITY VALUES

The following equations are used for calculating sheet resistance and resistivity values from dual-configuration four-probe measurements.

$$R_{s} = \frac{V}{I} K_{a} F_{T} F(t/S) = X K_{a} F_{T} F(t/S)$$
 (1)

$$\rho = \frac{V}{I} K_a F_T t F(t/S) = X K_a F_T t F(t/S)$$
 (2)

where:

 R_s is the sheet resistance of the wafer, in ohms;

 ρ is the volume resistivity of the wafer, in ohm centimeters;

(V/I) is the first-configuration (ASTM F84) voltage-to-current ratio (also called R_a), in ohms:

t is the wafer thickness, in centimeters;

S is the average probe separation, in centimeters;

F(t/S) is a thickness-related scaling factor (near unity for t < 0.4 S);

 F_T is a correction from the temperature of measurement to a reference temperature (23 °C);

 K_a is a geometric scaling factor that is calculated from electrical data in the two configurations; and

X is a shorthand for the voltage-to-current ratio in the first configuration.

The equations are applied at each measurement site to the average of the voltage-to-current ratio for the forward and reverse currents.

From the theoretical development of the dual-configuration measurement, the scaling factor, K_a , is determined from a transcendental equation from reference [8], but a simplified calculation that is a highly accurate representation is given by the following quadratic equation, also from reference [8]

$$K_a = -14.696 + 25.173 \left(\frac{R_a}{R_b}\right) - 7.872 \left(\frac{R_a}{R_b}\right)^2,$$
 (3)

where R_a is the voltage-to-current ratio in the first electrical configuration, and R_b is the voltage-to-current ratio in the second electrical configuration.

4.1 Rewriting the Equation to Relate to Evaluation of Uncertainty

In the ISO formulation of uncertainty, the standard uncertainty is the square-root of the sum of variances of the components evaluated by Type A procedures and of those evaluated by Type B procedures. Those components (e.g., short-, intermediate-, and longer-term measurement system imprecision) that enter through the measurement data are evaluated by statistical analysis of the actual measurements, in units of resistivity, and give a Type A standard uncertainty directly in units of resistivity. Those that enter through one of the scaling or correction factors in eq (1) or eq (2) must be multiplied by an appropriate prefactor to give a Type B standard uncertainty in the same units. The development of these prefactors is most readily done through a propagation of variance formulation for the variance of resistivity, $\sigma^2(\rho)$, in terms of the variances of the quantities in eq (2). The variance of resistivity can then be expressed as:

$$\sigma^{2}(\rho) = (t^{2}F^{2}(t/S)) \left[F_{T}^{2}\sigma^{2}(\chi) + \chi^{2}\sigma^{2}(F_{T})\right] + \left(\frac{\rho}{tF(t/S)}\right)^{2} \left[F^{2}(t/S) \sigma^{2}(t) + t^{2}\sigma^{2}(F(t/S))\right], \quad (4)$$

where χ is the product $X K_a$.

All certification and control experiment data that are supplied for statistical analysis are in units of resistivity, corrected to a temperature of 23 °C, with dimensions of ohm centimeters. The statistical variations in these data are principally manifestations of variations in the measured electrical quantities: the first term in eq (4); and to a lesser extent, variations in the temperature of measurement and the associated temperature correction: the second term in eq (4). Since each of the wafers being analyzed has a fixed assigned thickness value, there is no statistical variation due to thickness: the last two terms in eq (4). The statistical analyses

look at total change in resistivity from all sources and are not partitioned into variability of voltage and current or temperature correction. The results of the statistical analyses give values of uncertainty, in ohm centimeters, from which a Type A variance, in ohm centimeters squared, is calculated and then summed with the Type B variance.

All terms in eq (4) need to be considered in ISO Type B analyses of uncertainty related to measurement scale calibration errors. All terms, as written, have dimensions of ohm centimeters squared, but it is convenient to rearrange the first square-bracketed term of the equation so that it shows the same explicit dependence on ρ^2 that can be seen for the second square-bracketed term. By multiplying numerator and denominator of the first term by χ^2/F_T^2 , the equation can be rewritten as:

$$\sigma^{2}(\rho) = \frac{\rho^{2}}{\chi^{2} F_{T}^{2}} \left[F_{T}^{2} \sigma^{2}(\chi) + \chi^{2} \sigma^{2} (F_{T}) \right] + \frac{\rho^{2}}{t^{2} F^{2}(t/S)} \left[F^{2}(t/S) \sigma^{2}(t) + t^{2} \sigma^{2} (F(t/S)) \right]. \quad (5)$$

It is useful to summarize the nominal values of the various terms that appear as part of prefactors in eq (5). All such terms are sufficiently constant from wafer to wafer that use of nominal values will suffice. Nominal wafer thickness is 0.628 cm; the thickness-related scaling term F(t/S) is dimensionless and is taken as unity for all wafers because of the values of t/S for the wafers being certified. The temperature correction factor, F_T , is dimensionless and is very close to unity, being no smaller than 0.985, nor larger than 1.005 for any SRM wafer. The term K_a is dimensionless and has a slightly different value for each line of data for each wafer, but the value is always close to 4.50.

In order to facilitate Type B evaluation of measurement uncertainties, it is helpful to split the preceding equation into separate variance terms that can be related to the background discussions of Type B standard uncertainty evaluations in Section 3. These terms deal with:

electrical measurements:

$$\frac{\rho^2}{\chi^2 F_T^2} \left[F_T^2 \ \sigma^2(\chi) \right] = \rho^2 \left(\frac{\sigma^2(V/I)}{(V/I)^2} + \frac{\sigma^2(K_a)}{K_a^2} \right), \tag{5a}$$

temperature measurements:

$$\frac{\rho^2}{\chi^2 F_T^2} \left[\chi^2 \ \sigma^2(F_T) \right] = \frac{\rho^2 \ \sigma^2(F_T)}{F_T^2} , \qquad (5b)$$

and thickness measurements:

$$\frac{\rho^2}{t^2 F^2(t/S)} [F^2(t/S) \sigma^2(t) + t^2 \sigma^2 (F(t/S))] = \rho^2 \left(\frac{\sigma^2(t)}{t^2} + \frac{\sigma^2(F(t/S))}{F^2(t/S)} \right).$$
 (5c)

Thus, following this rearrangement, each of the contributions reduces to the relative variance of a variable times the square of the resistivity.

5. SOURCES OF MEASUREMENT UNCERTAINTY - DETAILED DISCUSSION

5.1 Type A Evaluations of Components of Uncertainty

The contributions to uncertainty from sources discussed in this section are evaluated solely from certification and control experiment data taken at the time of certification for each of the resistivity levels. A variance is calculated for each of the Type A contributions. These variances are then combined in a root-sum-of-squares fashion to give a standard deviation from the combination of effects; this standard deviation is the Type A standard uncertainty. Data from SRM 2547, at $200 \ \Omega$ -cm, are used in Appendix 2 to illustrate the analysis procedures used. Abbreviated summaries from SRMs 2541, 2542, 2545, and 2546, which follow the same procedures, are given in Appendices 3 through 6 (some of these latter appendices also contain analysis details of a specific additional term which was not pertinent to the data analyzed in Appendix 2). The statistical reports in the appendices state the standard uncertainties for the resistivities at the wafer-center and for the 5 mm and 10 mm measurement circles. In Section 7, those values from Appendices 2 through 6 are summarized, the variances from Type A and Type B analyses are tabulated, and the combined variances, combined standard uncertainties, and expanded uncertainties are given for both sheet resistance and resistivity for each of the SRMs.

5.1.1 Short-term precision; repeatability

There is expected to be negligible effect from wafer nonuniformity on the six measurements at the wafer center; ideally, these measurements would all have the same value. The standard deviation of the six values is a measure of the repeatability, or short-term precision, under tightly controlled conditions. The repeatability is evaluated from data taken over periods so short that there should be no changes in measurement environment, or wear or damage effects on the wafers or the probe. The variability among the data being analyzed is a combination of two effects, both of which cause fluctuations in the voltage-to-current ratios, and as a result, in the calculated K_a scaling factor that is based on those ratios. (See eq (1).) The first of these effects is the scatter in the electrical data due to pure electrical or electronic sources such as variations in probe contact quality, power supply noise, or DVM noise; the second is scatter in electrical data due to small fluctuations in probe separation, from one site to the

next, that is not fully corrected for by use of the dual-configuration technique. These two effects are the primary mechanisms causing short-term data scatter; they cannot be separated functionally, and there is no need for doing so. They are accounted for in calculations of short-term standard deviation of resistivity. Typical values for the standard deviation of a set of six measurements at the center of a wafer have been found to range from 0.03 % to about 0.30 % for single-configuration data and about 0.02 % to 0.12 % for dual-configuration data. The actual values for standard deviation depend somewhat on the probe used and on the wafer resistivity level. One of the causes of the spread in the observed values is the small sample size (six measurements in the NIST certification procedure) for calculating the standard deviation. The short-term precision for the certification process at each resistivity level is estimated from a pooling of variances of the wafer-center data from the wafers being certified and similar data from the wafers in both types of control experiments. There are typically 1000 or more degrees of freedom to this pooled estimate, depending on the number of wafers in the batch being certified.

5.1.2 Intermediate and longer term precision; reproducibility of wafer-center average value

Assuming there is no significant change in the probe used for certification measurements and no change in the resistivity of any of the wafers due to the probing process, it should be possible to remeasure any of the wafers and obtain average values that fall within limits based on the short-term precision value. In fact, this generally is not found. The small excess variation is believed due to changes in the measurement environment, such as power-line variations and changes in the radiated noise in the laboratory environment, humidity changes, etc., that are not readily identified over the short time spans used to measure individual wafers. Data from the replicate measurements on the check-wafer and also from the initial and final control experiment wafers are analyzed for a day-to-day (run-to-run) random variation in the response of the certification probe that is in excess of the pooled short-term standard deviation. In addition, comparisons of preliminary and final control experiment data for each of the probes on each of the control wafers are used to estimate any additional longer-term variations that are characteristic of the entire measurement system and changes in the environment, not just of the certification probe or the check-wafer. Such contributions to uncertainty are termed "long-term variations" in the statistical analysis reports. There are typically 50 degrees of freedom in the determination of the day-to-day variability in the analysis of the control-wafer data and 20 or more degrees of freedom in such a determination from the check-wafer data. There are 5 degrees of freedom for the calculation of long-term variability in the comparison of initial and final control experiment data. The same sets of data are also analyzed for possible systematic data trends in the measurement process or specimens and corrections terms applied or additional uncertainty components evaluated, as necessary. Such a systematic trend was identified for the 200 Ω cm wafers. It is discussed separately in Section 6, and the analysis of a resulting asymmetric modification of the uncertainty interval is given in Appendix 2.

5.1.3 Uncertainty due to the selection of a particular probe

It has been found, based on the analysis of many experiments, that resistivity measurement values obtained by four-point probe on bulk wafers have a small dependence on the probe being used. Experience at NIST shows this to be the largest residual error when ASTM Method F84 is used for measurement; measuring the geometric separation of probe impressions made on a polished wafer, as required by ASTM F84, does not adequately describe their functional electrical separation. This dependence is significantly reduced, but not eliminated, by use of the dual-configuration procedure. This may be thought of as an issue of the accuracy of the basic model of dual-configuration four-probe measurements applied to real probes having finite size contact areas and contacts that are not purely ohmic, but affected by metal-semiconductor interface effects. The result is a probe-dependent bias in the measured wafer resistance value that might normally be considered a systematic effect, the value of which could be evaluated, or estimated for any given probe. Because there is no model of the physics that causes the offset for a given probe, an estimate of the probable distribution of probe offset values cannot be done on a theoretical basis. A numerical evaluation of such a distribution could be done if given a sufficiently large number of probes, and the bias of a given probe could then be determined using the many-probe average as a point of reference. However, there are only five probe heads available for use in the certification procedure, thus making it impossible to obtain data from a sufficient variety of probe heads to generate a distribution of probe-dependence values.

Instead, the five available probe heads are treated as a random sample from the universe of probe heads, and sufficient replication data are taken with each probe head during the initial and final control experiments (Sec. 1.4) that a statistical estimate can be made for a variance term due to probes as a variable. Thus, while the choice-of-probe effect is most simply conceptualized as a systematic error, it is actually evaluated from statistical analysis of these replicate measurements as a Type A contributor to the standard uncertainty of certification. The initial and final control experiments incorporate data from both choices for wiring the second probe configuration, while the certification measurements use only one of those two choices. The initial and final control experiments are also analyzed for possible contribution to certification uncertainty due to small differences between the two choices for second configuration wiring.

5.2 Type B Evaluations of Components of Uncertainty

No corrections were applied to the SRM certification measurements for possible errors in voltage, current, or thickness values. However, a correction was applied for the difference between the temperature scale of the thermistor used to monitor measurement temperature and that of a precision mercury bulb thermometer which is the customary reference to a NIST-traceable temperature scale following the procedure of ASTM F84.

In this section, with one exception, a single value is calculated for uncertainty in electrical and thickness scales which is applicable to all resistivity levels. That exception is at 0.01 Ω cm, for which the value related to electrical measurements is almost twice as large as that for the

worst case from any of the higher resistivity levels. A separate value is given for 0.01 Ω cm. For the temperature correction term, it is necessary to calculate a separate value for each resistivity level.

In the remainder of Section 5.2, individual effects are considered significant, and are retained, if they are at least 0.01 % (one part in ten thousand) of the measured value. Values smaller than that are considered negligible.

5.2.1 Discussion of components related to electrical measurements

Measurement of Specimen Current — Four separate precision-current supplies are available, each calibrated and tested annually for ripple and noise. Measurement accuracy does not rely on this calibration, however. Instead, the measurement current is fed through a precision standard resistor in series with the wafer, and the voltage drop across the resistor is measured with the same 6-1/2 digit DVM (Hewlett-Packard model 3456) used for the silicon wafer measurements. Voltage measurements are taken with a resolution of 0.1 µV. Five precision resistors from 0.01 Ω to 1000 Ω are available. Each is calibrated periodically at NIST. The resistors have calibration uncertainties of 3 $\mu\Omega/\Omega$ to 5 $\mu\Omega/\Omega$. There is no meaningful change of value of these resistors due to temperature variations for the temperature excursions encountered in the lab. Standard resistor and wafer voltages are measured on the same range setting of the DVM. In typical practice, a standard resistor is selected for use so that it gives a voltage drop that is a factor of 1 to 10 times that of the specimen being measured; e.g., a 10 Ω standard resistor is used for the measurement of a 1 Ω cm wafer. Voltages measured across the standard resistor are typically 25 mV, generally stable to 1 µV and read to 0.1 µV. One of the two available solid-state power supplies is preferred for measurement because of the convenience of six-digit current selection; however, the regulation specifications for these current supplies (as a percent of full-scale) become marginal for the low currents used when measuring 100 Ω cm and 200 Ω cm SRMs, and it has been found preferable to switch to a vacuum-tube supply to maximize measurement current stability for these resistivities. Specifications for the current supplies and for the digital voltmeter are given in Table 1.

Measurement of Specimen Voltages — ASTM Method F84 requires that the measurement current be set to give a specimen voltage drop, between the two inner probes, of 10 mV to 20 mV. NIST measurements for SRMs 1521 to 1523 were taken in the restricted range of 10 mV to 12 mV. For the 100 mm SRMs, 2541 to 2547, reported here, measurements are taken in the still more restrictive range of 9.95 mV to 10.05 mV. (See Sec. 2.2.) (With a 1.59 mm probe point separation, this gives a maximum field of less than 7 mV/mm across the wafer.) Once the current is adjusted to give a specimen voltage in this range for the ASTM wiring configuration of the very first measurement at the wafer center, the power supply is left at this setting for all other measurements on that wafer. This specimen voltage range results in an acceptable number of digits of measurement resolution with minimal risk of Joule heating or minority carrier injection. The only exception to this procedure occurs for wafers with low resistivity (below about 0.05 Ω cm for a nominal 625 μm thickness) where use of a current supply having a typical 100 mA maximum output will result in a maximum obtainable specimen voltage that is below the range stated above. For the lowest resistivity

SRM, $0.01~\Omega$ cm, the specimen voltage at 100~mA is about 3.1~mV; this causes a somewhat larger relative uncertainty in the scale of the electrical measurements at this SRM level.

Typical stability of wafer voltage readings, as seen from the DVM display, ranges from $\pm 1~\mu V$ to $\pm 3~\mu V$ (depending upon resistivity, probe, and environmental conditions). In practice, after setting the switches for each desired voltage to be measured, the operator verifies that there is no drift in the DVM display for that setting by observing five to ten readings, and then causes the next DVM reading to be stored in the computer with the expectation that the scatter noted above represents a random error in the stored value.

Although the single DVM reading that is stored for each voltage or current measurement can be said to be in error as long as there is any scatter in the DVM displays observed by the operator, it is not necessary to do a first-principles propagation of error based on typical voltage scatter and eq (5a) in order to determine the random uncertainty in the voltage-to-current ratio. The standard deviation of a set of measurements taken in a fixed region of the wafer (e.g., the wafer center where material nonuniformity effects are negligible) encompasses the uncertainty due to digital voltmeter noise just described, as well as that due to variations in probe separation and probe contact quality. Thus, these sources of error are part of the short-term Type A uncertainty of measurement discussed in Section 5.1.1. It is not necessary to do any other analysis for these factors. Accuracy, or systematic error, of the digital voltmeter is limited by the 24 count, or 2.4 μ V specification. However, relative accuracy of the ratio measurement is better than 2.4 μ V and is essentially controlled by the accuracy of the standard resistor values. The effect of digital voltmeter accuracy on measurement uncertainty is given in Section 5.2.2.

General Integrity of the Electronic Instrumentation — This is basically a problem of elimination/rejection of noise, whether from electronic or thermal sources. When the current supplies are sent for calibration, they are also checked to verify that they are within the manufacturers' specifications for ripple and noise; see Table 1. The primary switch-matrix in the instrumentation utilizes heavy copper contact posts and twin seven-wiper blade construction designed to be thermal-voltage free. The common-mode and normal-mode noise rejection specifications for the DVM are stated for the case of a 1000 Ω measurement load; this value is exceeded, however, for all SRMs above 1 Ω cm. To test the effectiveness of noise rejection, as well as possible leakage currents, analog boxes with very large series resistors (that represent probe contact resistance, see ASTM F84) are measured with, and without, the series resistors in the circuit. This is done as a part of the preparation for certification of each SRM level. Worst-case experience shows that analog boxes simulating 10 000 Ω cm silicon experience a measurement difference (error) of about 0.20 % between these two setups. This decreases to about 0.02 % when simulating 1000 Ω cm silicon and is negligible for the simulation of 200 Ω cm and lower resistivity silicon.

Table 1. Manufacturers' Specifications for the Current Supplies and DVM Used for Certification

ELECTRONIC MEASUREMENTS Inc. Model C612 Constant-Current Supply*

OUTPUT RANGES:

1 μ A, 2.2 μ A, 5 μ A, multiplier x1, x10, x100 etc.

to 100 mA max. (0 to 100 % vernier each range)

STABILITY:

0.3 % of range setting (fixed line, load, and temp.)

CURRENT REGULATION:

0.1 % for 100 V step in compliance voltage

RIPPLE and NOISE:

0.04 % rms of range setting + 0.5 μA (negative ground)

0.04 % rms of range setting + $0.1 \mu A$ (positive ground)

(floating output is used for certification)

OUTPUT IMPEDANCE:

30 000 M Ω @ 1 μA to 500 k Ω @ 100 mA

ELECTRONIC DEVELOPMENT CORPORATION Model CR103 Constant-Current Supply

OUTPUT RANGES:

10 mA and 100 mA full scale; 6 digit setability

STABILITY (non-additive):

1 h 0.001 % of range

8 h 0.005 % "

1 Yr. 0.01 % "

RIPPLE and NOISE:

 $(0.1 \text{ Hz to } 100 \text{ kHz}) < 0.5 \mu\text{A}$

OUTPUT CONDUCTANCE:

 $0.1 \mu S$

TEMPERATURE COEFFICIENT: 0.0005 %/K

^{*}This current supply is operated at 50 %, or greater, of range setting.

Table 1. (cont'd.)

HEWLETT-PACKARD #3456 DVM

(All values are stated for the 100 mV range)

RESOLUTION (Least Count):

0.1 μV

INPUT IMPEDANCE:

 $>10^{10} \Omega$

MEASUREMENT ACCURACY:

For auto-zero on, filter off and ≥ 10 power cycle cycles): 24 h @ (23 ± 1) °C: $\pm (0.0022 \% \text{ rdg.} + 24 \text{ counts})$ 90 day @ (23 ± 5) °C: $\pm (0.0034 \% \text{ rdg.} + 24 \text{ counts})$

TEMPERATURE COEFFICIENT: ±(0.0002 % rdg. + 0.2 counts/°C)

NOISE REJECTION:

Normal mode, ac: 60 dB (1 kΩ max. Unbalance in low) Common mode, ac: 150 dB Common mode, dc: 140 dB

5.2.2 Evaluation of uncertainty in electrical measurement scale

Electrical measurement scale contributions to the variance of resistivity value are found from examining the right-hand side of eq (5a)

$$\rho^{2} \left[\frac{\sigma^{2}(V/I)^{2}}{(V/I)^{2}} + \frac{\sigma^{2}(K_{a})}{(K_{a})^{2}} \right]. \tag{6}$$

Ignoring temporarily the term in K_a , and replacing the current, I, with the ratio of standard resistor voltage to standard resistor value, V_s/R_s , in the first term, results in

$$\rho^{2} \left[\frac{\sigma^{2}(VR_{s}/V_{s})}{(VR_{s}/V_{s})^{2}} \right] = \rho^{2} \left[\frac{\sigma^{2}(V/V_{s})}{(V/V_{s})^{2}} + \frac{\sigma^{2}(R_{s})}{R_{s}^{2}} \right].$$
 (7)

Rather than expanding the term in (V/V_s) to get separate terms in $\sigma^2(V)$ and $\sigma^2(V_s)$, it is preferable to look at the way in which electrical measurement error affects the ratio, V/V_s , as a whole. It is assumed that the 2.4 μ V error (due to voltmeter accuracy limit statement) affects the measurements of V and V_s equally (both voltages are measured on the same meter, and in quick succession). Then the worst-case error in their ratio occurs when V_s is the largest multiple of V. It can be seen from Table 2 that this occurs when V_s approximately equals $3 \times V$.

For the resistivities above 0.01 Ω cm, the wafer voltage-drop is 10 mV, the standard resistor voltage-drop is 30 mV, and the worst-case ratio V/V_s , with no error in voltage values, is 0.333 333 3. A 2.4 μ V error in both V and V_s for these wafers causes a change in the ratio to 0.333 386 7. The difference of the two ratios is 0.000 053 4 and will be taken as a limit of error in the voltage ratios due to DVM least-count error. Squaring this value, and dividing by 3 (assuming a rectangular error distribution) gives a variance of 9.50×10^{-10} . The denominator, $(V/V_s)^2$, equals 0.111, so the contribution to variance from the first term above is: $(9.50 \times 10^{-10}/0.111) \rho^2$, or $8.56 \times 10^{-9} \rho^2$.

At 0.01 Ω cm, because of smaller measurement voltage levels, the contribution to uncertainty from electrical measurements is actually larger than the worst-case value for the SRMs above 0.01 Ω cm. For this SRM level, the ratio without voltage measurement error, is 3.1 mV/10 mV, or 0.310 000; and with a 2.4 μ V error, it is 0.310 166. The resulting error in the V/V_s ratio is 0.000 166. Squaring this, and dividing by 3, as above, gives 9.18 \times 10⁻⁹. The denominator, $(V/V_s)^2$, at 0.01 Ω cm, is 0.096. The resulting contribution to variance of resistivity, at 0.01 Ω cm, is 9.56 \times 10⁻⁸ ρ^2 .

The second term in eq (7),

$$\frac{\rho^2 \sigma^2(R_s)}{R_s^2} ,$$

can be shown to be negligible. The calibration uncertainty of all standard resistors used for the SRMs is $<5 \times 10^{-6}$ times the value of the resistor. Assuming a rectangular distribution for standard resistor calibration error, $\sigma^2(R_s)/(R_s)^2 = (2.5 \times 10^{-11}/3) = 8.3 \times 10^{-12}$. The contribution to variance related to standard resistor calibration error is 8.3×10^{-12} ρ^2 and is negligible. Likewise, possible contributions due to drift, or to temperature dependence of standard resistor values are negligible compared to the one part in ten-thousand criterion noted above.

Table 2. Standard Resistor Values, and Typical Measurement Voltages for Each of the SRM Levels

Nominal SRM Value (Ω cm)	Standard Resistor (Ω)	SRM Wafer Voltage, <i>V</i> (mV)	Std. Res. Voltage, V_s (mV)
0.01	0.1	3.1	10
0.1	1	10	25
1	10	10	25
10	100	10	25
25	100	10	11.5
100	1000	10	30
200	1000	10	14

The other contribution to uncertainty due to electrical measurement scale error comes from the second term in eq (5a)

$$\rho^2 \left[\frac{\sigma^2(K_a)}{K_a^2} \right]$$

 K_a has the following characteristics. It is the solution to a transcendental equation based on two configurations of electrical data taken at each measurement site. The solution has been approximated by a quadratic equation in the argument R_a/R_b , where R_a is the ratio of V/I in the first (ASTM) wiring configurations and X_b is the V/I ratio in one of the two choices for the second configuration. Specifically, the quadratic equation is

$$K_a = -14.696 + 25.173 \left(\frac{R_a}{R_b}\right) - 7.872 \left(\frac{R_a}{R_b}\right)^2$$

The accuracy of the fit over the range $1.20 < R_a/R_b < 1.32$ is reported to be better than 0.05 % [9]. For the wafer diameter, measurement locations, and probe size used in this SRM certification, the ratio, R_a/R_b , is approximately 1.255. There are small variations, from about 1.25 to 1.26, which encompass both the effects of electrical measurement noise and small fluctuations in the separation of adjacent pairs of probe pins from one measurement position to the next. A ratio of R_a/R_b of 1.255 results in a K_a value of about 4.50. Over this restricted range, the accuracy of fit of the quadratic, is actually about 0.01 %.

There are two independent considerations in evaluating σ^2 (K_a). The first is the relative inaccuracy, 0.01 %, of the quadratic representation of the transcendental equation. With the assumption of a uniform probability distribution, it results in a contribution to the variance of $[\{(0.000\ 1\ K_a)^2/3\}/K_a^2] \times \rho^2$, or $3.33 \times 10^{-9}\ \rho^2$.

The second is the error in K_a that would occur because of an error in measured voltages. For a nominal value of $R_a/R_b=1.255$, and any of the SRMs above 0.01 Ω cm, a voltage measurement error of 2.4 μ V would cause an error in R_a/R_b of no more than 0.000 07. This causes a change (error) in K_a of about 0.000 38. Again assuming a rectangular distribution of error, this means that the voltage error contribution to variance is $[\{(0.000\ 38)^2/3\}/(4.50)^2] \times \rho^2$ or about $2.39 \times 10^{-9}\ \rho^2$. For the 0.01 Ω cm SRM, and under the same assumptions, a voltage error of 2.4 μ V causes an error in the ratio, R_a/R_b , of 0.000 24. This, in turn, results in an error in K_a of 0.001 29 and a contribution to the variance of ρ^2 at 0.01 Ω cm of $[\{(0.001\ 29)^2/3\}/(4.50)^2] \times \rho^2$, or about $2.74 \times 10^{-8}\ \rho^2$.

Adding these terms to that for possible error due to the quadratic representation of the transcendental equation, the variance in ρ^2 due to possible error in the factor, K_a , is $3.07 \times 10^{-8} \rho^2$ at $0.01 \ \Omega$ cm, and is $5.72 \times 10^{-9} \rho^2$ for SRMs above $0.01 \ \Omega$ cm.

No specific additional systematic error terms due to instrumentation integrity have been identified in the resistivity range of these SRMs other than the 0.02 % offset that has been seen with the 1000 Ω analog box. Noise, due to poor contact quality, radiated signal pickup, or other sources, may be present. It is believed to contribute scatter, in the low microvolt level, to the data, and show up as a component of the standard deviation of the data. It is possible, but has not proven necessary, to integrate measurements on the DVM for 100 power-line cycles, instead of the customary 10 cycles, to suppress the effects of ac pickup.

Therefore, the total contribution to variance of resistivity due to electrical measurement considerations discussed above is 1.263×10^{-7} ρ^2 at $0.01~\Omega$ cm and 1.428×10^{-8} ρ^2 at all higher resistivities.

5.2.3 Evaluation of uncertainty components related to temperature measurements

The variance of resistivity value due to temperature measurement errors arises as follows: During resistivity measurement, each wafer is placed on a copper block which is both massive, to maintain temperature stability, and made of a good thermal conductor, to enhance the speed of equilibration of temperature between the surface where the wafer is located and the block's interior where the thermistor temperature sensor is located. A thin mica film provides electrical insulation between the wafer and the copper block. The measured temperature (maintained in the range 22 °C to 24 °C for all SRM wafer measurements and observed to be stable to 0.1 °C, or better, for any given SRM wafer) is used in conjunction with an empirically evaluated temperature coefficient of resistivity for silicon to correct the measured resistivity to the standard value of 23 °C. The temperature coefficient of resistivity for silicon, which is a function of both resistivity and conductivity type, was evaluated at NBS in the mid-1960's. This temperature coefficient is used internationally and is part of a

standard measurement procedure (ASTM F84) for silicon resistivity near room temperature. It is expected that all users of these SRMs for application to silicon technology will use the same temperature coefficients for interpretation of their "unknown" or "test" wafers. No evaluation of uncertainty of the coefficient itself is made here.

The thermistor was calibrated against a precision mercury bulb thermometer over the range $15~^{\circ}\text{C}$ to $35~^{\circ}\text{C}$. The mercury bulb thermometer itself was calibrated by NIST with a stated uncertainty of $\pm 0.03~^{\circ}\text{C}$, or better. Thermistor resolution is better than $0.01~^{\circ}\text{C}$. Transfer uncertainty between glass bulb and thermistor is estimated to be no worse than $0.02~^{\circ}\text{C}$; a value of $0.02~^{\circ}\text{C}$ will be used. The largest potential error is that the copper block temperature may not be the same as that of the wafer. This could be due to warm or cool air currents from the room ventilation system affecting the wafer and block exterior. Tests of consistency of resistivity measurement with controlled temperature increase and decrease indicate that potential error between sensor and wafer is less than $\pm 0.08~^{\circ}\text{C}$. The calibrations of the glass bulb thermometer and that of the thermistor are added to give a worst-case temperature calibration error of $\pm 0.05~^{\circ}\text{C}$. This is added linearly to the possible wafer-sensor offset of $0.08~^{\circ}\text{C}$ to give a worst case total temperature error of $0.13~^{\circ}\text{C}$.

Because all possible temperature errors were added linearly to calculate worst-case error, above, it is overly conservative to assume a uniform distribution of error to calculate a variance, and a triangular distribution for the temperature error is assumed instead. Thus, the variance of the distribution of possible temperature error is $(0.13 \text{ °C})^2/6 = 0.002 \text{ 82 (°C)}^2$.

To minimize possible temperature error in practice, wafers are kept in the vicinity of the measurement station for at least 24 h prior to measurement, and have at least 1 min to stabilize on the copper block before taking measurements. Possible errors in resistivity values due to temperature enter through the term from the right-hand side of eq (5b)

$$\rho^2 \; \frac{\sigma^2(F_T)}{F_T^2} \; ,$$

where the temperature correction of resistivity, F_T , has the form,

$$F_T = 1 - C_T (T - 23 \, {}^{\circ}C)$$
,

where C_T is the temperature coefficient of resistivity, in degree Celsius⁻¹, and T is the temperature at which measurements are made, in degree Celsius.

The variance in F_T is really the variance in temperature (given above) times the square of the temperature coefficient, $(C_T)^2$. Since the coefficient, C_T , varies noticeably as a function of

resistivity value, Table 3 summarizes the values of the temperature coefficient used in the calculation of uncertainty.

Table 3. Temperature Coefficients of Resistivity of Silicon for the Nominal Values of the SRMs

Nom. Res. (Ω·cm)	0.01	0.1	1	10	25	100	200
Temp. Coeff. (°C) ⁻¹	0.0031	0.0041	0.0071	0.0082	0.0083	0.0083	0.0083

^{*}Exact values are given on the certificate for each SRM wafer.

The variance of resistivity value due to temperature error is 0.002 82 $(C_T)^2 \rho^2$. Because of the difference in the values of C_T for the various SRM levels, the variance in resistivity value due to temperature error is given in Table 4.

Table 4. Variance in Resistivity Value Due to Temperature Error

SRM Resist Ω·cm	ivity $\sigma^2(F_T)$	Contribution to Variance of Resistivity
0.01 0.1 1 10 25 100 200	2.71×10^{-8} 4.51×10^{-8} 1.42×10^{-7} 1.90×10^{-7} 1.94×10^{-7} 1.94×10^{-7} 1.94×10^{-7}	$2.71 \times 10^{-8} \rho^{2}$ $4.51 \times 10^{-8} \rho^{2}$ $1.42 \times 10^{-7} \rho^{2}$ $1.90 \times 10^{-7} \rho^{2}$ $1.94 \times 10^{-7} \rho^{2}$ $1.94 \times 10^{-7} \rho^{2}$ $1.94 \times 10^{-7} \rho^{2}$ $1.94 \times 10^{-7} \rho^{2}$

5.2.4 Evaluation of the uncertainty components related to geometry measurements

In single configuration (ASTM F84) measurements by four-point probe, it is necessary to measure accurately the wafer diameter, the wafer thickness, the average separation between the probe pins, and variability thereof, in order to calculate geometry-related scaling factors that convert measured voltage/current ratios to sheet resistance and resistivity values. In dual-configuration measurements, only the measurement of wafer thickness and the average probe separation (for thicker wafers) enters into the calculation of sheet resistance and

resistivity. The following discussion deals with errors in geometry measurements as they relate to possible uncertainties in the certification values.

Comments on Nonideality of a Lapped Surface — A lapped surface texture is used to optimize electrical stability of the SRMs and to improve contact quality between the probe and wafer. The wafer thickness, in centimeters, is used to multiply sheet resistance values to convert them to resistivity values. Fractional errors in thickness values are reflected 1:1 as fractional errors in calculated resistivity values. The lapped wafer surface has a peak-and-valley texture that is related to, but generally smaller in size than, the abrasive used to do the lapping. Even though the lapping process used to prepare the SRM wafers is known to give total (macroscopic) thickness uniformity better than obtainable on as-cut or polished wafers, the existence of the surface texture precludes there being a unique thickness value at any location on the wafer. (The 100 mm SRM wafers for SRMs 2541 to 2547 were lapped with a simultaneous two-side lapping process. An abrasive grit size of about 12 µm was used for the four lowest resistivities and a $7 \mu m$ grit size for the three highest resistivities. The earlier 50.8 mm (2 in) diameter NBS silicon resistivity SRMs utilized a one-side-at-a-time process and a 5 µm abrasive. As a result, the 100 mm wafers have much improved macroscopic thickness uniformity, but a somewhat coarser surface texture compared with earlier SRMs.)

Measured thickness values are somewhat dependent on the method of measurement. Electromechanical-, capacitive-, acoustic-, or air-gauges are not expected to respond the same to the hills and valleys of a textured surface or to average over the same surface area. A mechanical method that measures front-surface-to-back-surface peak-to-peak thickness is the most idealized conceptually when dealing with these circumstances, and was used for thickness measurements of the SRM wafers. However, the peak heights on a lapped surface are somewhat variable on both wafer faces (resulting in small local fluctuations in peak-to-peak thickness and some sensitivity to the location where the thickness measurements are made). Figure 3 attempts to illustrate the situation of defining and measuring thickness on a textured surface using an electromechanical gauge.

Calibration and Control of the Electronic-Micrometer — Wafer thicknesses of the SRMs were measured with an electronic-micrometer having a resolution of 0.05 μ m and a short-term repeatability of about 0.1 μ m. The instrument's specifications state that its accuracy is $\pm 0.1~\mu$ m if the ambient temperature is kept at 20 °C ± 1 °C. The requirement of a temperature of 20 °C is based on the temperature at which the instrument was calibrated by the manufacturer. While the laboratory at NIST in which the instrument is used maintains the required 1 °C temperature stability, the nominal working temperature is typically 23 °C. To maintain the calibration accuracy of the micrometer, standard practice is to calibrate, and to recheck, the instrument a number of times a day against precision gauge blocks traceable to NIST and having thicknesses that are comparable to the SRM wafers. The gauge readout was reset, as necessary, to match the gauge block value. Thus, the thickness measurement for the SRMs was a process of transfer of thickness value from a gauge block through the thickness

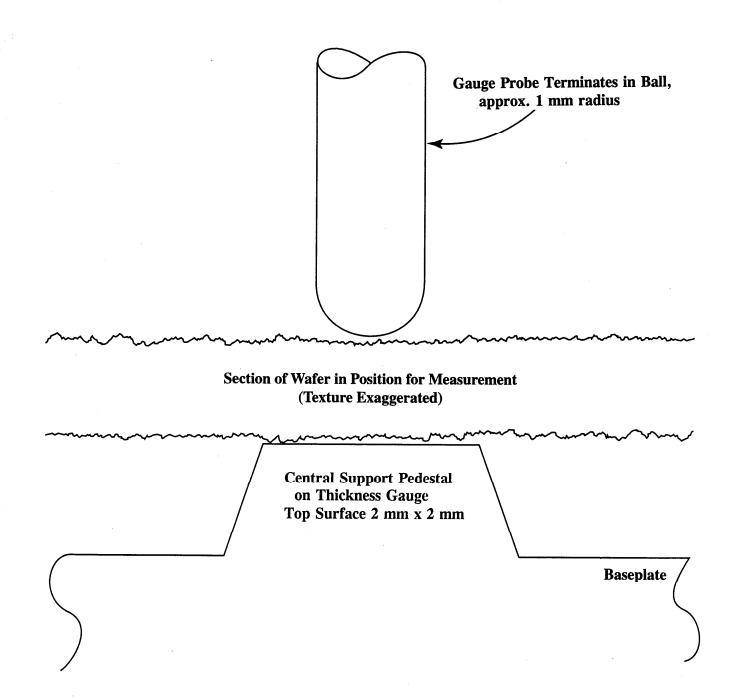


Figure 3. Conceptual drawing of wafer with two textured surfaces during thickness measurement by electromechanical gauge.

gauge to the wafer. Several error components can be identified that will affect the accuracy of this transfer. The thickness gauge specifications are given in Table 5.

[Note: These resistivity SRMs are not intended to serve as thickness calibration standards, and that it goes beyond the scope of this work to be able to relate the performances of electromechanical-, capacitive-, acoustic-, air-gauge, and other thickness methodologies on lapped surface wafers.]

Table 5. Specifications for Haidenhain Certo 60 Thickness Measurement Instrument

Measurement Resolution	0.05 μm
Measurement Accuracy	±0.1 μm
(At an Operating Temp of 19 °C to	21 °C)
Wafer Backside-Reference Pedestal	2 mm x 2 mm
Probe Tip Radius	1 mm
Probe Working Force	1 N
•	

5.2.5 Evaluation of uncertainty due to thickness measurement scale

Possible errors in the thickness measurement scale contribute to uncertainty of resistivity directly through the first term in eq (5c),

$$\rho^2 \frac{\sigma(t)^2}{t^2} .$$

Based on the average SRM wafer thickness of 0.0628 cm, the denominator has a value of $0.003~94~\text{cm}^2$.

Two error mechanisms contribute to a Type B estimate of variance of thickness values. The first relates to the calibration of the thickness measurement tool with precision gauge blocks. Three blocks with thicknesses of 0.024 in, 0.025 in, and 0.026 in (0.060 96 cm, 0.063 50 cm, and 0.066 04 cm), i.e., just spanning all expected values of wafer thickness, and having NIST-traceable thicknesses known to better than 0.000 004 in (0.000 01 cm) are used. In the tool calibration procedure, the tool is adjusted to read the known thickness of the 0.025 in (0.06350 cm) block and required to read the other two within 0.15 μ m (0.000 015 cm) of their stated calibration values. This is 50 % larger than the uncertainty of individual gauge block calibration values. The value 0.000 015 cm is taken as the half-width of the rectangular

distribution of possible error assignable to the calibration of the thickness measurement tool. As a result, there is a contribution to variance of resistivity of $1.90 \times 10^{-8} \rho^2$ from possible tool calibration error.

The second error mechanism relates to the transfer of the thickness measurement scale to the SRM silicon wafers. Various tests of consistency of wafer thickness values suggests that a rectangular distribution with a 0.1 μ m half-width should be sufficient to account for the thickness transfer error term. This results in a contribution to variance of resistivity of $8.37 \times 10^{-9} \, \rho^2$. Combining these two terms gives a variance of resistivity value directly due to variance of thickness of $2.74 \times 10^{-8} \, \rho^2$.

5.2.6 Evaluation of uncertainty due to thickness/probe separation scaling factor A scaling factor, F(t/S), is used to correct the calculated sheet resistance values for layers of finite thickness (greater than about 0.4 times the average probe spacing). Error in either the wafer thickness or in average probe separation value contributes to the variance of sheet resistance or resistivity through the second term in eq (5c),

$$\rho^2 \frac{\sigma^2 (F(t/S))}{F^2(t/S)} ,$$

where the denominator is effectively unity.

The scaling factor, F(t/S), for dual-configuration measurements is similar to that, F(w/S), for single-configuration measurements in that they both asymptotically approach unity for values of w/S just below 0.4. These scaling factors are virtually identical for values of the ratio, w/S, below about 0.45, but diverge noticeably in value for wafer thicknesses that are a large fraction, or a multiple, of the probe separation.

ASTM F84 recommends the simplification that this factor be set to unity when the ratio of wafer thickness to average probe separation is 0.4 or less. For all larger values of the ratio, the scaling factor is then computed from summation of a specified series and takes on values decreasing from unity as the ratio increases above 0.4. The ratio, 0.4, exactly corresponds to a wafer thickness of 635 µm (0.025 in) and a probe separation of 1587 µm (0.0625 in). This is the nominal separation of the probes being used for SRM certification, and the SRM wafers were, in fact, purchased with a target thickness of 625 µm. Some fraction of the wafers in a given SRM batch will exceed the ratio, 0.4, if only by a small amount, simply due to fabrication process tolerances. For the seven SRM levels, a total of 34 wafers (out of approximately 800) had thicknesses such that the t/S ratio exceeded 0.4; the worst-case value of the ratio was 0.4008.

Examination of the scaling factor shows that it actually has a value of 0.9995, not unity, for thickness-to-probe-spacing ratios that are infinitesimally above 0.4. When the procedure for

certification of these SRMs was devised, it was not known exactly how much variation in wafer-to-wafer thickness would be encountered. To avoid the inconsistency in scaling factor that would result from using a default value of unity for ratios up to 0.4, and then a calculated value of the scaling factor for all higher values of the ratio t/S, a decision was made to calculate and apply a correction term for all values of the thickness/probe separation ratio. The result is to improve the SRM wafer-to-wafer consistency for resistivity value as a function of thickness, but to introduce an offset for most SRM wafers that makes their stated resistivity 0.04 % to 0.05 % smaller than if the asymptotic value of unity had been used for this scaling factor. The exact amount of the offset for a given SRM wafer can be found, if needed, by comparing to unity the value of this scaling factor as printed on the certificate for that wafer. This offset is incorporated in both sheet resistance and resistivity values. There is no error, or uncertainty, term developed to relate to this change from the procedure of ASTM F84.

To calculate the variance in the scaling factor due to uncertainty in the measurements of thickness and probe separation, typical results for probe separation measurements and thickness data from one of the SRM levels are used. Following the procedures of ASTM F84, probe separations can be measured to a resolution of about 1 μ m and with a typical precision for 10 readings of about 0.06 % (1 μ m).

Wafer thickness and probe-spacing values for the 25 Ω cm SRM level are used to calculate the variance of the wafer-thickness probe-spacing scaling factor. For this SRM level, the slightly larger upper end wafer thicknesses relative to the spacing of the probe used make the sensitivity of this term a little larger than for the other SRM levels. For this SRM, assuming no error in thickness or probe separation value, the ratio, t/S, ranges from 0.387 26 (F(t/S) = 0.999 632), to 0.400 83 (F(t/S) = 0.999 506). A worst-case combination of probe separation error (0.0002 cm assumed) and wafer thickness error (0.000 025 cm assumed) causes a change in the scaling factor value of about 0.000 03 (a relative change of 0.003 %). Using this value as a half-width (the error could also be the same amount in the opposite direction), and assuming a rectangular distribution, the variance of F(t/s) is 5.07×10^{-10} . Thus, this contribution is negligible.

6. UNANTICIPATED EFFECTS

During the course of certification of the seven SRMs, two effects were encountered that had not been experienced previously and which were thus partially, or wholly, outside the design of the control experiments. The first of these was a shift, or drift, in measured resistivity during the first few rounds of probe measurements on the 200 Ω cm SRMs. The second was a sensitivity of the measured resistivity to background illumination level for the 1 Ω cm (SRM 2543) and the first batch of 10 Ω cm (SRM 2544) wafers.

6.1 Resistivity Shift with Repeated Probing

The phenomenon of resistivity shift with repeated probing is documented in Sections 2.1 and 3.3 of the analysis of SRM 2547 which is given in Appendix 2. It shows up as a decrease of measured resistivity with successive sets of probe measurements made within a period of days or weeks. It was found to occur for some but not all wafers tested, and where it exists, it is stronger for some probes than for others. The shift is not totally cumulative, but appears to saturate.

Additional measurements of the original control wafers more than a year after the acquisition of the certification data showed nearly the same effect as shown in Appendix 2. The first of these additional measurements started at almost the same value as originally (i.e., an upward recovery of value had occurred in the interim), followed by a gradual decrease in resistivity by about the same amount as previously, then reached an asymptotic value. As previously noted, some of the control wafers suffered the effect; others did not. The observed shift did not accumulate beyond a few tenths of a percent.

The mechanism for this shift is unknown. It is not believed to be experienced with repeated eddy current measurements, but this hypothesis was not tested. It is expected that when wafers from SRM 2547 are first measured by the user, they will manifest the resistivity (sheet resistance) values listed on the certificate, and if measured by four-point probe, some of them will show small decreases of resistivity if replicate probe measurements are made within a period of days, or perhaps weeks. The additional term added to the estimated uncertainty interval due to analysis of this effect on the control wafers is believed to fully cover any manifestation of this effect to the user.

6.2 Photosensitivity of Resistivity Value

Measurements being made separately from the certification of these SRMs, and after the time when most of the SRMs had been measured for certification, showed that certain types of silicon had a resistivity value that was dependent on the level of background illumination. Extensive previous experience with four-probe measurements of many silicon specimens, particularly the types used for previous SRMs, had shown that normal laboratory-level fluorescent illumination had no observable effect on the measurement value. It had been seen that high resistivity silicon (perhaps $1000~\Omega$ cm, and higher) must be measured in the dark. It was also seen that bright incandescent illumination, with a significant component of penetrating infrared radiation, would inject hole-electron pairs that would decrease the measured resistivity with a very rapid recovery (because of short minority-carrier lifetimes) to higher values when that illumination was turned off.

Previous resistivity SRMs up to 200 Ω cm, fabricated from float-zone or neutron-transmutation-doped silicon, showed no sensitivity of resistivity to normal laboratory levels of fluorescent lighting, and certification measurements were taken on them without a dark-box enclosure. The current 100 mm diameter SRM wafers were fabricated from boron-doped

Czochralski silicon from $0.01~\Omega$ cm to $10~\Omega$ cm and from float-zone grown, NTD-doped silicon for the highest three resistivity levels. These choices of silicon types were made specifically to optimize within-wafer uniformity of resistivity for the various SRM levels.

The photosensitivity of resistivity that was detected subsequent to SRM certification occurred on boron-doped Cz silicon wafers that were not related to the SRM wafers. Subsequent testing showed that the effect was measurable from a few tenths of an ohm centimeter to the highest resistivity boron-doped Cz silicon obtainable, approximately 80 Ω·cm. The magnitude of the shift in resistivity was found: 1) to be as high as 2.5 %, 2) to depend roughly on resistivity level, and 3) to be present on all boron-doped Cz silicon wafers available, independent of supplier and wafer surface type. Auxiliary tests, on wafers from the 1 Ω cm and 10 Ω cm SRM crystals, also showed a correlation between the magnitude of the effect and the interstitial oxygen level. The effect could not be detected at all on boron-doped float-zone silicon or on any phosphorus-doped silicon. Tests were then made of the existence and magnitude of this effect on wafers from the four boron-doped Cz silicon crystals that had already been certified for SRMs. No effect could be detected for the 0.01 Ω cm or $0.1~\Omega$ cm resistivity levels. A photoeffect as large as 0.4~% and decreasing to about 0.15~%. as a function of wafer position in the starting crystal, was detected for wafers from SRM 2543, at 1 Ω cm. The effect ranged from 0.6 % to 1.2 % for wafers from the crystal initially used for SRM 2544, at 10 Ω cm.

The photosensitivity is unusual in its very long decay time from lower resistivity in normal room illumination to higher resistivity in the dark. Typical times for decay to the asymptotic value typical of the new illumination state ranged from about 2 min to more than 20 min. Wafers used for SRMs 2543 and 2544 were at the lower end of this time scale.

Because of the significantly large value of the photoeffect for wafers from the original $10~\Omega$ cm boron-doped Cz crystal, these wafers were invalidated for use as SRMs. It was possible to purchase a sufficient quantity of $10~\Omega$ cm wafers grown by the float-zone process and phosphorus-doped by the NTD technique to be able to retain the $10~\Omega$ cm SRM level using these replacement wafers. The NTD wafers are nearly as uniform as the boron-doped Cz silicon wafers they replaced and are suitable for use as SRMs since they show no evidence of a photosensitivity. The complete set of certification and control measurements have been completed on the NTD wafers. At the time of publication, analysis of those data is not complete.

The case for the 1 Ω cm SRM level was not so straightforward. It was not possible to get float-zone grown, NTD-doped wafers that are irradiated heavily enough to produce 1 Ω cm silicon. Possible replacement Cz silicon wafers doped with phosphorus were expected to be free of photosensitivity, but to have sufficiently large nonuniformity of resistivity as to be unacceptable for use as standards. No other alternative could be identified, and a choice had to be made between voiding the 1 Ω cm SRM level altogether and a judicious use of the 1 Ω cm wafers already measured. The decision was made to retain only the best of the original 1 Ω cm SRM wafers, i.e., those wafers having the lowest amount of photosensitivity,

about 0.25 % and below. This will allow the retention of about half of the originally certified batch of 125 wafers. The task of selection was made easy because the supplier for those wafers laser-engraved a unique serial number on each wafer in the sequence the wafers were taken from the saw. The magnitude of the photoeffect had been found to decrease monotonically from the low numbered toward the high numbered wafers.

During an additional analysis, yet to be completed, of the certification data and auxiliary data, an estimate will be made of a new component of uncertainty, due to the state of illumination. This estimate will be based on measurements at normal laboratory illumination levels, measurements in the dark, and measurements at noticeably higher than normal levels of illumination. The latter is to simulate possible shifts to lower values of resistivity that might occur in user facilities that have higher illumination levels than were present during certification. Because this analysis is not complete, there are certain omissions for the $1~\Omega$ cm level in the summary tables of uncertainty values in Section 7. When completed, the analysis will be documented in a supplement to this report. All measurement control procedures, ISO Type B components of uncertainty, and the general analysis procedures for ISO Type A components detailed in this report remain valid for this SRM.

Two notes of caution are in order regarding the use of moderate to lightly boron-doped Cz silicon wafers, regardless of source, for resistivity standards. Both are based on the assumption that photosensitivity, of the type described here, is a universal characteristic of boron-doped Cz silicon. First, it is not sufficient, in general, simply to take the certifying data in darkened surroundings. Any user of such a standard who is not able to take measurements in similarly darkened surroundings will experience a different resistivity value, and the difference between the dark-level and illuminated-level values may not be characterized adequately. Second, because the decay time for the photosensitivity is so long, it is relatively easy, using most commercial, automated instrumentation, to be fooled about whether a photosensitivity exists for a given wafer. Only a series of measurements over a period of minutes is likely to reveal the drift that is caused by this photosensitivity. There is a related consideration, for a wafer certified in the dark, that will be measured in a darkened, or shrouded, user-instrument, but which has been stored in illuminated surroundings. Such a wafer will have to be allowed to equilibrate with the darkened interior of the instrument for a number of minutes before valid readings can be taken.

7. COMPILATION OF UNCERTAINTY COMPONENTS

This section summarizes the Type A standard uncertainty terms for resistivity from Appendices 2 through 6 and the Type B variance terms from Section 5. It uses these inputs to obtain the combined variance, u_c^2 , the combined standard uncertainty, u_c , and the expanded uncertainty for each of the following parameters. The expanded uncertainty, U_c , is stated on the SRM certificates for: 1) average resistivity at the wafer center; 2) average sheet resistance at the wafer center; and 3) individual sheet resistance measurements at locations on the 5 mm and 10 mm circles.

The values of Type A standard uncertainty in the appendices are given only for resistivity values. To convert these to values appropriate to sheet resistance, it is necessary only to divide them by the average SRM wafer thickness, 0.0628 cm. Separate values are needed for average sheet resistance at the wafer center and for individual measurements on the 5 mm and 10 mm radius circles. There are three considerations for converting the values of Type B variance of resistivity, given in Section 5, to values of variance of sheet resistance. First, since sheet resistance values do not depend on wafer thickness, only the terms in Section 5 from the variance of the electrical and temperature measurements contribute to Type B variance of sheet resistance. Second, Sections 5.2.2 and 5.2.3 give the contributions of electrical and temperature measurement variations to the variance of resistivity; it is necessary to divide those variance-of-resistivity terms by the square of the average SRM wafer thickness, (0.0628 cm)², to scale to the variance of sheet resistance. Third, the Type B variance terms are estimates of measurement scale error, and are the same for average measurements at the wafer centers and for individual measurements on the two small circles.

7.1 Summary of Statistical Analysis Parameters from the Appendices

This section summarizes the information given in Appendices 2 through 6. It gives the symbols used in the statistical analyses, the components of Type A standard uncertainty that they represent, and a table of values obtained for these components for five of the SRM levels. It also gives a summary of the Type A standard uncertainty values for wafer center averages and for individual values on the 5 mm and 10 mm radius circles for both resistivity and sheet resistance.

Table 6. Components Identified in Statistical Analyses of Certification and Control Experiment Data

s_{ϵ}	Short-term imprecision of certification probe
s_{δ}	Run-to-run measurement variability
S_{γ}	Longer-term measurement variability
$s_{c}^{'}$	Uncertainty of non-zero correction for certification probe bias
s_{Δ}	Uncertainty of non-zero correction for probing induced drift, wafer-probe interaction
S_{cfig}	Uncertainty of non-zero correction for probe configuration difference
$\frac{s_{\text{cfig}}}{a/\sqrt{3}}$	Uncertainty of correction for probe configuration
	(where the best correction = 0); [Type A estimate, but based on limit of error]
<i>b</i> /√3	Uncertainty of correction for probe bias
	(where the best correction = 0); [Type A estimate, but based on limit of error]
and	
Type	A standard uncertainty, $u_i = (s_{\epsilon}^2/(n) + s_{\delta}^2 + s_{\gamma}^2 + s_{c}^2 + s_{\Delta}^2 + s_{cfig}^2)^{1/2}$, where $n = 1$ for individual measurements on the circles, and $n = 6$ for the
	where $n = 1$ for individual measurements on the circles, and $n = 6$ for the
	average value at the center.

Table 7. Values of the Components Identified in Statistical Analyses for the Various SRMs in Appendices 2 through 6, $m\Omega \cdot cm^{*\#}$

SRM	$s_{ m c}$	s_{δ}	s_{γ}	Se	s_{Δ}	$s_{ m efig}$	a/√3	<i>b</i> /√3
2541	0.001 83	0.001 04	4 0.004 00	0	0	0	0	0.000 47
2542	0.062	0.032	0.004	0	0.011	0	0	0.016
2545	14.14	3.31	3.01	0	0	0	2.89	0
2546	72.	13.4	14.6	5.1	0	0	0	0
2547	138.	64.	129.	5.	10.	0	0	0

^{*}For ease of reading, this table is expressed in terms of milliohm centimeters. *Statistical analyses of SRMs 2543 and 2544 are not complete.

Table 8. Type A Standard Uncertainty Values, u_i , Taken from Appendices 2 to 6*

SRM	RESISTIVITY at center $(m\Omega \cdot cm)$	RESISTIVITY on circles (mΩ·cm)	SHEET RESISTANCE at center (mΩ)	SHEET RESISTANCE on circles $(m\Omega)$
2541	±0.004 23	±0.0045	±0.0673	±0.0716
2542	±0.045	± 0.072	±0.725	±1.16
2545	±7.8	±15.1	±125.	±241.
2546	±35.8	± 74.8	±570.	±1190.
2547	±155.	±199.	±2470.	±3180.

^{*}Statistical analyses of SRMs 2543 and 2544 are not complete.

7.2 Type A and Type B Variance Terms

In this section, a table of Type A variance terms is constructed from the squares of the Type A standard uncertainty values given in Table 8. A table of Type B variance values, obtained from the analyses in Section 5, is also given. Finally, a table of the combined variance, u_c^2 obtained by adding the Type A and Type B variances, is then given.

Table 9. Type A Variance Values, u_i^2 , Obtained by Squaring the Entries in Table 8*

			SHEET	SHEET
CD) (RESISTIVITY	RESISTIVITY	RESISTANCE	RESISTANCE
SRM	at center	on circles	at center	on circles
	$(\Omega \mathrm{cm})^2$	$(\Omega \cdot \text{cm})^2$	(Ω^2)	(Ω^2)
	11	11	0	. 0
2541	1.79×10^{-11}	2.02×10^{-11}	4.53×10^{-9}	5.13×10^{-9}
2542	2.02×10^{-9}	5.18×10^{-9}	5.26×10^{-7}	1.35×10^{-6}
2545	6.08×10^{-5}	2.28×10^{-4}	1.56×10^{-2}	5.81×10^{-2}
2546	1.28×10^{-3}	5.59×10^{-3}	3.25×10^{-1}	1.42
2547	2.40×10^{-2}	3.96×10^{-2}	6.10	10.1

^{*}Statistical analyses of SRMs 2543 and 2544 are not complete.

Table 10. Type B Variance Values, u_j^2 , Calculated from Summation of Terms in Sections 5.2.1 through 5.2.4

SRM	RESISTIVITY at center & on circles $(\Omega \cdot cm)^2$	SHEET RESISTANCE at center & on circles (Ω^2)
2541 2542 2543 2544 2545	2.51×10^{-11} 8.68×10^{-10} 1.84×10^{-7} 2.32×10^{-5} 1.47×10^{-4}	5.68×10^{-9} 1.51×10^{-7} 3.96×10^{-5} 5.18×10^{-3} 3.30×10^{-2}
2546 2547	2.36×10^{-3} 9.43×10^{-3}	5.28×10^{-1} 2.11

Table 11. Combined Variance Values, u_c^2 , from Addition of Terms from Tables 9 and 10*

SRM	RESISTIVITY at center $(\Omega \text{ cm})^2$	RESISTIVITY on circles $(\Omega \cdot \text{cm})^2$	SHEET RESISTANCE at center (Ω^2)	SHEET RESISTANCE on circles (Ω^2)
2541	4.30×10^{-11}	4.53×10^{-11}	1.02×10^{-8}	1.08×10^{-8}
2542	2.89×10^{-9}	6.05×10^{-9}	6.77×10^{-7}	1.50×10^{-6}
2545	2.08×10^{-4}	3.75×10^{-4}	4.86×10^{-2}	9.11×10^{-2}
2546	3.64×10^{-3}	7.95×10^{-3}	8.53×10^{-1}	1.95
2547	3.35×10^{-2}	4.90×10^{-2}	8.21	12.2

^{*}Statistical analyses of SRMs 2543 and 2544 are not complete.

7.3 Combined Standard Uncertainty and Expanded Uncertainty.

This section gives values of the combined standard uncertainty, u_c , and the expanded uncertainty, U, based on a coverage factor k = 2. The combined standard uncertainty values are the square roots of the entries in Table 11 for the combined variance.

Table 12. Combined Standard Uncertainty Values, u_c^*

SRM	RESISTIVITY at center (Ω·cm)	RESISTIVITY on circles (Ω·cm)	SHEET RESISTANCE at center (Ω)	SHEET RESISTANCE on circles (Ω)
2541	±0.000 006 56	±0.000 006 73	±0.000 101	±0.000 104
2542	±0.000 053 8	±0.000 077 8	±0.000 823	±0.001 22
2545	±0.014 4	±0.019 4	±0.220	± 0.302
2546	±0.060 3	±0.089 2	±0.924	±1.39
2547	±0.183	±0.221	±2.86	±3.49

^{*}Statistical analyses of SRMs 2543 and 2544 are not complete.

Table 13. Expanded Uncertainty Values, U (Coverage factor k = 2)*

SRM	RESISTIVITY at center (Ω·cm)	RESISTIVITY on circles (Ω·cm)	SHEET RESISTANCE at center (Ω)	SHEET RESISTANCE on circles (Ω)
2541	±0.000 013 1	±0.000 013 5	±0.000 202	±0.000 208
2542	±0.000 108	±0.000 156	±0.001 65	±0.002 45
2545	±0.028 8	±0.038 7	±0.441	± 0.604
2546	±0.121	±0.178	±1.85	±2.78
2547 [#]	-0.498, +0.366	-0.575, +0.443	-7.83, +5.73	-9.08, +6.98

^{*}Statistical analyses of SRMs 2543 and 2544 are not complete.

7.4 Corrections Applied to Measured Values

This section summarizes bias corrections that must be made to measurement results, as acquired, because of effects that were identified during statistical analyses of the SRM certification experiment data. These corrections are explained in the appropriate appendices. They are given for resistivity and sheet resistance values. All resistivity and sheet resistance values shown on the SRM certificates have already been corrected for these bias terms.

Table 14. Bias Corrections Applied to Measured Values

SRM	AMOUNT	SOURCE
2541	Subtract 0.000 000 472 Ω cm (0.000 007 52 Ω) [This is a negligible amount]) Wiring Configuration Bias
2542	Subtract 0.000 037 5 Ω·cm (0.000 597 Ω)	Wiring Configuration Bias
2543		Wiring Configuration Bias and
		Photoeffect Terms to be Evaluated
2544	_	Wiring Configuration Bias Detected
2545	None	
2546	Add 0.0393 Ω·cm (0.626 Ω)	Probe Bias
2547	Subtract 0.0490 Ω cm (0.78 Ω)	Probe Bias
	(wafer drift term of $0.132 \Omega cm$ is built into ex	panded uncertainty)

[#]Asymmetry is due to a contribution of 1.32 Ω cm, 2.10 Ω , from wafer-probe interaction, i.e., to a drift in value.

8. CONCLUSION

At the time the certification procedure for these SRMs was being developed, the quantitative design objective was to support the goal for layer resistivity stated in the SEMATECH Mega-IC Workshop, i.e., measurements with a 1 % accuracy and a 0.5 % repeatability. Prior to the evaluation of an uncertainty component for the photosensitivity effect of the 1 Ω cm SRM, 2543, the uncertainty values listed in Table 13 can be used to assess how well the objective was met. Table 13 shows that the relative expanded uncertainty for center resistivity average measurement is 0.13 %, or less, for four of the SRMs listed and is less than 0.22 % for the 200 Ω cm SRM, even after allowing for the short-term drift caused by repeated probing. The results for SRM 2544 at 10 Ω cm, not yet finalized, appear to be close to 0.15 % for center point resistivity average. The uncertainty values at the center are smaller yet if stated for sheet resistance average values. The relative uncertainty for individual sheet resistance values on either of the measurement circles is 0.17 %, or less, for the SRMs listed at, or below, 100 Ω cm, and is 0.25 % for the 200 Ω cm SRM. Thus, these SRMs should serve quite well to support the SEMATECH-stated goals.

9. ACKNOWLEDGMENTS

It is a happy privilege to acknowledge the dedication and persistence of Donnie Ricks throughout all the preliminary testing, the certification, and follow-up testing that was necessary for the certification of these SRMs. Without her conscientious attention to detail, the uncertainty levels noted in the report could not have been achieved. Thanks also to Mike Thomas for his able assistance throughout the certification. A special thank you goes to Jane Walters for her diligent and meticulous work in preparing this report from a number of disparate-format pieces of source material.

REFERENCES

- [1] ASTM Method F84-93, "Standard Method for Measuring Resistivity of Silicon Wafers with an In-Line Four-Point Probe," Annual Book of ASTM Standards Vol. 10.05, West Conshohocken, PA 19428.
- [2] Recommendations for Specific Standards Actions, Proceedings of the SEMATECH Workshop on Silicon Materials for Mega-IC Applications, New Orleans, Louisiana, January 30-31, 1991, p. 9.
- [3] van der Pauw, L. J., A Method of Measuring Specific Resistivity and Hall Effect of Discs of Arbitrary Shape, Phil. Res. Rep. 13, 1-9 (1958).
- [4] Guide to the Expression of Uncertainty in Measurement, ISBN 92-67-10188-9, 1st Ed. ISO, Geneva, Switzerland, (1993).

- [5] Taylor, B.N. and Kuyatt, C. E., Guidelines for Evaluating and Expressing the Uncertainty of NIST Measurement Results, NIST Technical Note 1297, U.S. Government Printing Office, Washington DC (1994).
- [6] ASTM Method F1529-96, "Standard Method for Sheet Resistance Uniformity Evaluation by In-Line Four-Point Probe with the Dual-Configuration Procedure," Annual Book of ASTM Standards Vol. 10.05, West Conshohocken, PA 19428.
- [7] Harris, F. K., Electrical Measurements (John Wiley & Sons, New York, 1952), p. 431.
- [8] Perloff, D. S., Gan, J. N. and Wahl, F. E., Dose Accuracy and Doping Uniformity of Ion Implant Equipment, Solid State Technol. 24 (2), 112-120 (1981).
- [9] Perloff, D. S., Four-Point Probe Correction Factors for Use in Measuring Large Diameter Doped Semiconductor Wafers, J. Electrochem Soc. 123 (11), 1745-1750 (1977).

Appendix 1. Summary of Important SRM Wafer Material and Measurement Condition Parameters

This appendix consists of two tables that summarize important useful silicon wafer characteristics and electrical measurement conditions that apply to the various resistivity levels of SRMs 2541 to 2547. Table 1 lists the nominal resistivity, crystallographic orientation of the SRM wafer surfaces, crystal growth type, dopant species, and the commercial supplier for the wafers for each of the SRMs. Table 2 lists the four-point probe identification, the serial number and nominal resistance of the standard resistor used, as well as the nominal value of the measurement current used for certification of these same SRMs.

Table 1. Silicon Wafer Characteristics That Apply to Various Resistivity Levels of the SRMs 2541 to 2547

ier
con Corp.
con Corp.
con Corp.
er Siltronic
l Semi. A/S
l Semi. A/S
l Semi. A/S

Table 2. Electrical Measurement Conditions That Apply to Various Resistivity Levels of the SRMs 2541 to 2547

SRM	Probe	Standard Resistor Nominal Value/ Serial Number	Nominal Measurement Current
2541	283	0.1 Ω / 1771494	100 mA
2542	281	1Ω / 1594503	28 mA
2543	283	10Ω / 1593079	2.8 mA
2544	283	100Ω / 1598893	260 μΑ
2545	2062	$100 \ \Omega$ / 1598893	110 μΑ
2546	2362	$1000 \ \Omega$ / 1592167	29 μΑ
2547	SRM1	$1000 \ \Omega$ / 1592167	14 μΑ

Appendix 2. Analysis of Certification Data and Control Experiments for SRM 2547

1. GENERAL COMMENTS AND SUMMARY OF TYPE A STANDARD UNCERTAINTY COMPONENTS

1.1 Introduction

This appendix documents the statistical analysis leading to the certification of wafers from crystal 21566 for SRM 2547 and outlines a general procedure for analysis of other SRMs in the series 2541 through 2547. The results of the analyses of the remaining SRMs are briefly summarized in the following appendices. In addition to the three random components and the first three systematic components listed below which are common to all SRMs, this report also treats a small drift effect that was not found with any of the other SRMs.

The 137 wafers in this issue have nominal resistivities of $200 \,\Omega$ cm, and the wafers are assumed to be identical with regard to wafer face. Certification measurements are made with a single probe, identified as SRM1. Data consist of measurements at six locations on each of three circles located at 0 mm, 5 mm, and 10 mm from the center of each wafer, with the wafer face chosen at random with respect to the crystal growth direction. Sources of error which could contribute to the uncertainties of the certified values and which are examined in this appendix are: probe imprecision, run-to-run variability, long-term variability, differences between wiring configurations, differences between wafer faces, differences among probes (probe SRM1 bias), and wafer drift with probing, which was an unanticipated effect.

Only the standard deviation associated with probe spacing and electronic imprecision can be estimated from the certification data for the SRM wafers. A series of control experiments was carried out to identify and estimate error components which cannot be addressed by the certification measurements.

Measurements on a check-standard, chosen at random from the wafers in the issue, were made routinely during the certification procedure to: identify any anomalous behavior, document the stability of the process, and estimate a day-to-day component of measurement error. For this issue, the check standard is wafer #150; it was measured only with the certification probe SRM1.

Pre- and post-certification control experiments with five probes on five wafers with both second-configurations, b1 and b2, were repeated on 6 days. These measurements are

intended to estimate both the random and systematic components of the measurement process. The next section summarizes the Type A standard uncertainty for SRM 2547. It also gives a statement of how the uncorrected term due to wafer drift contributes to the expanded uncertainty. Tables 1 and 2 in Section 1.2 give an executive summary of the terms that contribute to the Type A standard uncertainty. The details of the calculation of the component terms are given in subsequent sections.

1.2 Certified Resistivities and Uncertainties

The average of six measurements on the 0 mm circle of each wafer, corrected for the effect of probe SRM1, is reported as the certified resistivity value. The Type A standard uncertainty associated with the certified value for the wafer center is

$$u_i = \left(s_c^2 + s_{\gamma}^2 + s_{\delta}^2 + \frac{1}{6} s_{\epsilon}^2 + s_{\Delta}^2\right)^{1/2} = 0.155 \ \Omega cm.$$

The expanded uncertainty (coverage factor k = 2) allows for an uncorrected systematic error of $-\Delta$. See Section 3.3 for details. Because the uncorrected systematic error is always in one direction, the expanded uncertainty interval is nonsymmetric and is expressed as

Certified value -
$$(2 u_i + \Delta)$$
, Certified value + $2 u_i$,

where $2 u_i = 0.310 \ \Omega \cdot \text{cm}$, and $\Delta = 0.132 \ \Omega \cdot \text{cm}$.

Individual measurements on the 5 mm and 10 mm circles for each wafer, corrected for the effect of probe SRM1, are reported as certified values on the certificates. The Type A standard uncertainty associated with each of these individual certified values is

$$u_i = (s_c^2 + s_v^2 + s_b^2 + s_e^2 + s_\Delta^2)^{1/2} = 0.20 \ \Omega \cdot cm$$
.

The expanded uncertainty interval for individual measurements is then expressed by

Certified value -
$$(2 u_i + \Delta)$$
, Certified value + $2 u_i$

where 2 u_i for individual measurements = 0.40 Ω ·cm.

Table 1. Components of Type A Standard Uncertainty for Crystal 21566, SRM 2547 with Probe SRM1, Ω·cm

Type	Source	Std dev	Estimate
Random	Imprecision of probe SRM1	3S.	0.138
Random	Run-to-run measurement variability	$S_{\mathcal{S}}$	0.064
Random	Long-term measurement variability	\hat{s}^{\prime}	0.129
Random	Standard deviation of correction for probe SRM1 bias*	S	0.005
Random	Standard deviation of wafer drift with probing	S_{Δ}	0.010
Systematic Systematic Systematic	Difference between configurations b1 and b2 Difference between front and back faces of wafer Neglected correction for wafer drift with probing	◁	Negligible Negligible 0.132

*A correction of -0.049 Ω ·cm due to probe SRM1 bias is applied to wafer center resistivity data for all wafers.

Table 2. Sources of Variation for Crystal 21566, SRM 2547 with Probe SRM1, Ω·cm

Source of error	Experiment	RMSE ^a	DF	Relationship
	SRM certifications	0.1374	685	
Probe imprecision	Pre-certification	0.1586	150	S
	Post-certification	0.1134	150	
	Check standard	0.1367	105	
		-		
	Pooled	0.138	1090	
Run-to-run	Pre-certification	0.0857	20	$(s_c^2 + \frac{1}{2}s_s^2)^{1/2}$
	Post-certification	0.0926	20	(3-0 0-)
	Check standard	0.0772	20	
	Pooled	0.0854	09	
	Pre- and post-certification	0.1116	5	$(s + \frac{1}{5}s + \frac{1}{5}s + \frac{1}{2\pi}s)$
Long-term	#900 wafers	0.1477	∞	(3-00 0-0 h)
	Pooled	0.1350	13	

associated with the individual effects, namely, imprecision, run-to-run variability, probe variability, and long-term variability, are computed ^aThe root-mean-square error, RMSE, estimates within each source-of-error category are pooled in the table above. Standard deviations using the relationship shown in the last column, with the results summarized in Table 1 of this Appendix.

^bDegree of Freedom.

^cThis column expresses the error components that comprise the pooled value in each "source of error" series of experiments; see reference at end of this Appendix.

2. RANDOM COMPONENTS

2.1 Pre- and Post-Certification Control Experiments

A nested experiment was performed with five probes on five wafers. Six measurements were made at the center position of each wafer with each probe; this sequence was repeated on 6 days; and the entire experiment was conducted twice, i.e., prior to and at the conclusion of the certification experiment. The temporal error model for one probe and one wafer is

$$y_{ijk} = \mu + \gamma_i + \delta_{ij} + \epsilon_{ijk} \qquad i = 1, 2; \ j = 1, ..., 6; \ k = 1, ..., 6 \eqno(1)$$

where μ is the average value, γ_i is a component for long-term error; δ_{ij} is a component of run-to-run measurement error; and ϵ_{ijk} represents short-term measurement imprecision error associated with the probe and electronics.

For this SRM, the pre- and post-certification measurements were made on opposite faces of the same wafers. Thus, there is a question as to whether the differences (see Fig. 1) between the pre- and post-certification measurements are caused by: (1) biases between faces; (2) drift on the wafer surfaces; or (3) long-term error in the measurement process.

- 1) Because the faces for the pre-certification experiment were chosen at random, it is unlikely that the differences, which are consistently in one direction, are caused by a front-to-back bias on the wafers. See Figure 1 where resistivity measurements on the five wafers are plotted versus the month/day of measurement. Also, measurements made 2 to 3 months after the conclusion of the certification process on additional wafers called #901, #902, #903, and #904 show differences which are consistently in the opposite direction.
- 2) There are not sufficient data from these experiments to judge inherent wafer drift.
- 3) The behavior of the pre- and post-certification data, which show strong correlations across wafers with time, is consistent with a components of variance model such as eq (1).

Sources of error and root-mean-square error terms (RMSE) for this model are in Table 2. For analysis of the initial and final control experiments, the first day's measurements were omitted. Estimates are made for each wafer individually and then pooled over wafers. The last column of the table shows the relationships between the results of the various experiments and the terms in the temporal error model above.

Crystal 21566

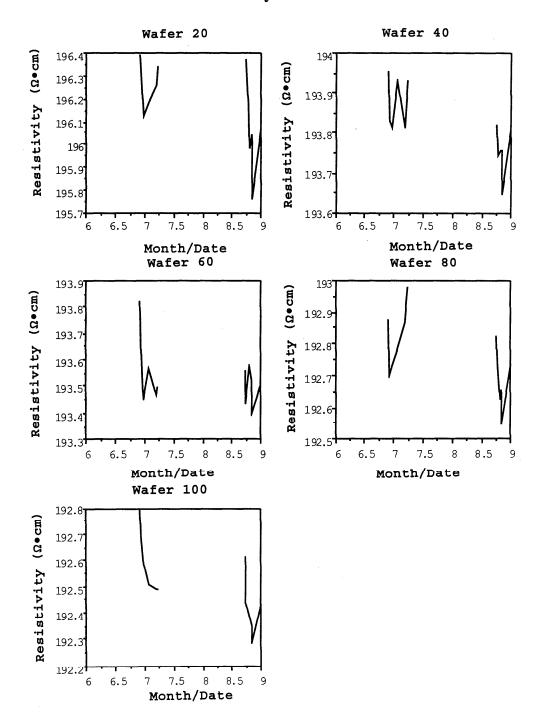


Figure 1. Resistivity (Ω ·cm) on five control-wafers from crystal 21566 with probe SRM1 plotted versus the month/day of measurement, showing change between and within pre- and post-certification experiments.

2.1.1 Precision of probes

The standard deviation, s_{ϵ} , is directly computed from six measurements at the center and estimates the precision for each probe. These standard deviations are shown in Table 3; the pooled values are also shown in Table 2.

Table 3. Within-Run Standard Deviations, s_{ϵ} , Pooled over Five Wafers and Six Runs, Ω cm

	Std. dev, s_{ε}		Std. dev, s_{ϵ}	
Probe	Config b1	DF	Config b2	DF
SRM1	0.1586	150	0.1907	150
281	0.2235	150	0.2468	150
Pre 283	0.2139	150	0.2389	150
2062	0.1645	150	0.2043	150
2362	0.1520	150	0.1635	150
CD164	0.1134	150	0.1200	150
SRM1	0.1134	150	0.1280	150
281	0.2102	150	0.2217	150
Post 283	0.1687	150	0.2115	150
2062	0.1568	150	0.1770	150
2362	0.1269	150	0.1374	150
Pooled				
Value: SRM1	0.1379	300	0.1624	300

2.1.2 Run-to-run measurement variability from pre- and post-certification control experiments

Standard deviations and averages computed from the six repetitions with each probe on each wafer are shown in Table 4. For this purpose, the first run with each probe on each wafer has been discarded. Each standard deviation is then estimated with four degrees of freedom. The pooled standard deviation for SRM1 of 0.089 20 Ω ·cm with 40 degrees of freedom incorporates both probe imprecision and day-to-day measurement error as shown in the relationship column of Table 2.

Table 4. Run-to-Run Component of Error, Crystal 21566 Averages and Standard Deviations for Last Five Runs on Each Control-Wafer, Ω·cm

		Pre-c	ertification	Post-ce	ertification
Wafer#	Probe	Average	Std dev	Average	Std dev
20	SRM1	196.2431	0.0875	196.0078	0.1574
40	SRM1	193.8663	0.0615	193.7433	0.0605
60	SRM1	193.5259	0.0795	193.4869	0.0693
80	SRM1	192.8096	0.1127	192.6597	0.0777
100	SRM1	192.5503	0.0835	192.3768	0.0596
20	281	196.2443	0.1380	196.0423	0.2617
40	281	193.8445	0.1105	193.7325	0.0948
60	281	193.5903	0.0935	193.4285	0.1399
80	281	192.7595	0.0826	192.6768	0.1939
100	281	192.5428	0.1189	192.3991	0.0930
20	283	196.1670	0.0937	195.9598	0.1525
40	283	193.7223	0.0499	193.6426	0.0951
60	283	193.4253	0.0536	193.3253	0.0992
80	283	192.7630	0.0396	192.5120	0.0945
100	283	192.4705	0.0545	192.3259	0.0824
20	2062	196.1481	0.1042	195.9211	0.2248
40	2062	193.8217	0.0957	193.7494	0.0711
60	2062	193.4647	0.0723	193.4411	0.0355
80	2062	192.7436	0.0727	192.6205	0.1538
100	2062	192.4263	0.0412	192.3818	0.0644
20	2362	196.1432	0.0884	195.8630	0.2282
40	2362	193.7696	0.0681	193.7181	0.0667
60	2362	193.4426	0.0581	193.3722	0.0775
80	2362	192.7206	0.0920	192.5816	0.0810
100	2362	192.4557	0.1279	192.2694	0.1589
Pooled					
Value:	SRM1		0.085 73		0.092 55
			/-		

2.1.3 Long-term measurement variability from the control-wafers

Averages for each wafer from the pre- and post-certification experiments are shown in Table 5. The differences are assumed to be the result of a long-term component of measurement error. The standard deviations as estimated from the pre- and post-certification averages represent probe imprecision, day-to-day error, and long-term measurement error as shown in the relationships column of Table 2.

Table 5. Pre- and Post-Certification Averages with Probe SRM1, Ω·cm

Wafer#	Pre-Certification	Post-Certification	Difference	Std dev	DF
20	196.2431	196.0078	-0.2353	0.1664	1
40	193.8663	193.7433	-0.1230	0.0870	1
60	193.5259	193.4869	-0.0390	0.0276	1
80	192.8096	192.6597	-0.1499	0.1060	1
100	192.5503	192.3768	-0.1735	0.1227	1
Pooled					
Value:				0.1116	5

2.1.4 Long-term measurement error from #900 series wafers

Averages of six center measurements made 2 to 3 months after the certification procedure are shown in Table 6. The measurements were made on a random selection of additional wafers numbered #901, #902, #903, and #904. The differences are assumed to be the result of a long-term component of measurement error. The standard deviations as estimated from the September and October averages represent probe imprecision, day-to-day error, and long-term measurement error as shown in the relationships column of Table 2. The fact that the differences shown in Table 5 for the control-wafers are always negative, whereas the differences observed for the #900 series of wafers are nearly always positive, is taken to indicate that this is not an inherent systematic effect. Therefore, no systematic correction term is applied.

Table 6. Long-Term Changes in Measurement Process with SRM1, Ω·cm

Wafer	Face	September	October	Difference	Std dev	DF
901	1	192.748	192.839	+0.091	0.0643	1
902	1	195.654	195.593	-0.061	0.0431	1
903	1	200.866	201.096	+0.230	0.1626	1
904	1	190.982	191.381	+0.399	0.2821	1
901	2	192.704	192.974	+0.270	0.1909	1
902	2	195.478	195.533	+0.055	0.0389	1
903	2	200.955	201.098	+0.143	0.1011	1
904	2	191.198	191.367	+0.169	0.1195	1
Pooled						
Value:					0.1477	8

2.2 Check-Standard Measurements

Twenty-three measurements (averages of six center measurements each) with probe SRM1 on wafer #150 were made over the course of the certification experiment. The initial drop in resistivity after the first day, which can be seen in Figure 2, is assumed to be the result of wafer-probing damage. The first day's measurements are omitted from the analysis. The slope of a straight line fit to the remaining 21 measurements as a function of time is not significant, indicating that the measurement process is not drifting. Therefore, only run-to-run variations in the measurement process and probe imprecision are reflected in the standard deviation which is shown in Table 2.

Crystal 21566

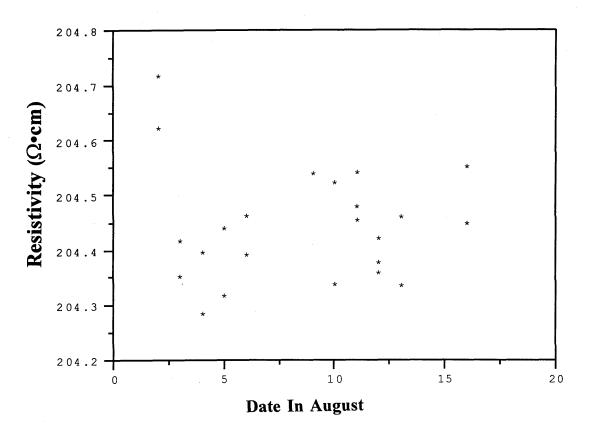


Figure 2. Resistivity measurements (Ω ·cm) on check-wafer #150, with probe SRM1 as a function of time (date in August)

3. SYSTEMATIC COMPONENTS

3.1 Systematic Differences between Probe-Wiring Configurations b1 and b2

In the pre- and post-certification experiments, six measurements at the center with the probe in configuration b1 were immediately followed by six measurements with the probe in configuration b2. The differences between configurations b1 and b2 for the pre- and post-certification measurements are shown in Figure 3. Averages and standard deviations for each probe over 6 days and five wafers are shown in Table 7. The t-statistic,

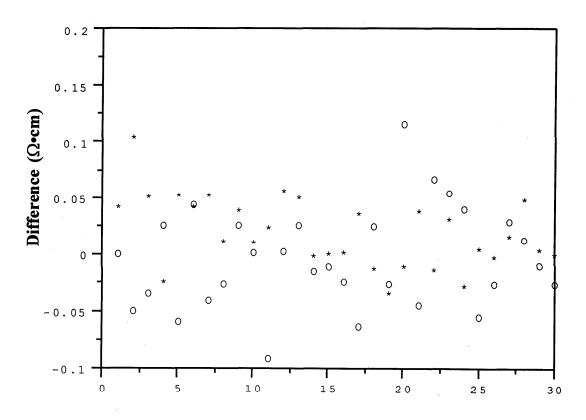
$t = \sqrt{30}$ Average/Std dev,

shows no evidence of a significant difference between configurations b1 and b2 for the pre-certification measurements and some evidence of a difference for the post-certification measurements. These differences for the post-certification measurements appear to be caused by the measurements on the first two wafers. No uncertainty from this source is assigned.

Table 7. Average Differences over All Control-Wafers between Configurations b1 and b2 for Probe SRM1, Ω·cm

	Pre-ce	rtification		Post-certification	
Probe	Average	Std dev	DF	Average Std dev I)F
SRM1	-0.002 44	0.044 49	29	0.020 56 0.031 38	29

Crystal 21566



Measurement Number for 5 Wafers over 6 Days Each

Figure 3. Differences between wiring configurations b1 and b2. Five wafers in random order over each of 6 days, with probe SRM1, Ω -cm.

Legend: O = pre-certification; * = post-certification

3.2 Differences among probes

The probes in the SRM certification are assumed to be a random sample of similar probes. However, certification using a single probe can have a systematic effect on the measurements. For this SRM, the measurements with SRM1 are found to be high relative to measurements with the other probes. Figures 4 and 5 show differences from the mean for each wafer plotted by probe. The systematic nature of these differences argues that the measurements made with SRM1 (identified by the number 1 in the plots) should be corrected to the average of the five probes based on the pre- and post-certification control measurements.

The estimated correction is calculated as the average of the differences in the table below to be $\hat{C}=-0.049~\Omega$ cm. The standard deviation of the differences is divided by $\sqrt{10}$ to obtain the standard deviation of the correction, $s_{c}=0.0050~\Omega$ cm. The correction, \hat{C} , is applied to all certified resistivity values, and its standard deviation is taken as a Type A component of uncertainty.

Table 8. Differences between Multi-Probe Average and Probe SRM1 for Each of the Control-Wafers, Ω·cm

Wafer	Pre-certification	Post-certification
20	-0.054	-0.049
40	-0.061	-0.026
60	-0.036	-0.076
80	-0.050	-0.050
100	-0.061	-0.026

Mean difference = $-0.049 \Omega \cdot \text{cm}$

Standard deviation, $s = 0.0159 \ \Omega \cdot cm$

Correction to be applied = $-0.049 \Omega \cdot \text{cm}$

Standard deviation of correction, $s/\sqrt{10} = 0.005 \ \Omega \cdot cm$

Crystal 21566

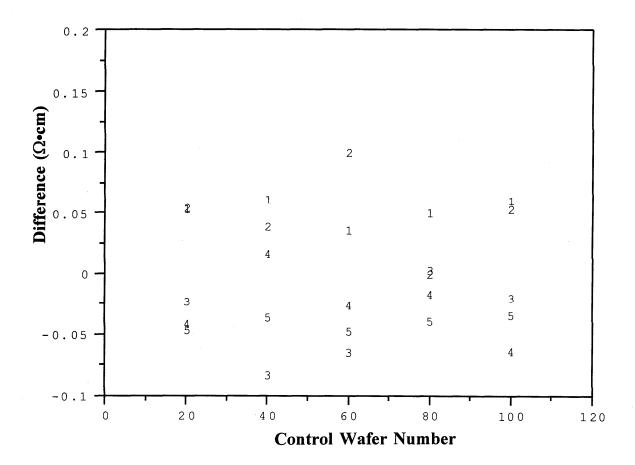


Figure 4. Differences between individual probe responses and multiprobe average, for pre-certification measurements on each of the control-wafers.

Plot symbol code: 1 = SRM1; 2 = 281; 3 = 283; 4 = 2062; 5 = 2362

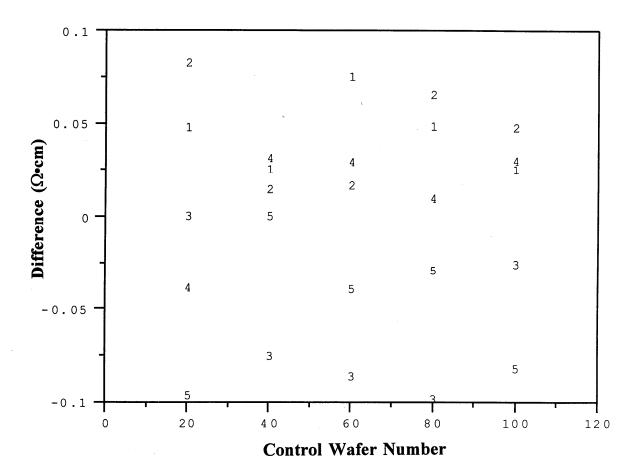


Figure 5. Differences between individual probe responses and multiprobe average, for post-certification measurements on each of the control-wafers.

Plot symbols: 1 = SRM1; 2 = 281; 3 = 283; 4 = 2062; 5 = 2362

3.3 Initial wafer damage

There is evidence from previous as well as present experiments that initial probing may change the surface characteristics of the 200 Ω ·cm wafers. The phenomenon is not totally understood nor always consistent, but has displayed itself as an initial drop in resistivity. The resistivity on check-wafer #150 dropped 0.3 Ω ·cm after the first day's measurements and then leveled off. For the pre- and post-certification measurements with SRM1, the resistivities always dropped after the first measurement for probe SRM1 (see Fig. 1). The

average drop is $\Delta = 0.132~\Omega$ cm, and the standard deviation is $s = 0.0427~\Omega$ cm. A correction of - $\Delta/2$ would assume an equal probability of initial damage between 0 and - $\Delta~\Omega$ cm. However, we choose to apply a correction for this asymmetry not to the data, but rather to the calculation of the uncertainty in Section 1.2. The term Δ is added to the lower limit of the expanded uncertainty. The standard deviation associated with Δ is $s/\sqrt{20}~\Omega$ cm or $0.010~\Omega$ cm, and is treated as a Type A component of uncertainty in the analysis.

Table 9. Drop in Resistivity between First and Second Measurements of Control-Wafers with Probe SRM1, Ω ·cm

Wafer	Pre-certification	Post-certification
20	0.086	0.185
40	0.122	0.072
60	0.174	0.122
80	0.177	0.093
100	0.113	0.176

The resistivity dropped after the first measurement (a measurement is the average of six readings at the wafer center) for all five control-wafers in both the pre- and post-certification experiments with probe SRM1, i.e., ten times out of ten possibilities. However, for the other probes, the number of times there was a drop after the first measurement in the same experiments was as follows: 2062 - seven out of ten possibilities; 281 - four out of ten; 2362 - four out of ten; and 283 - six out of ten possibilities. Thus, the effect is stronger for probe SRM1 than for any of the other probes. Plots of the complete data from the pre- and post-certification experiments are given in Figures 6 and 7.

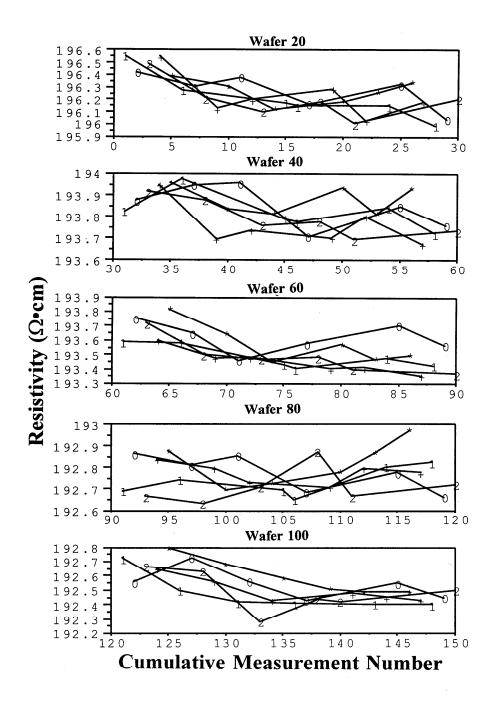


Figure 6. Resistivity (Ω ·cm) from pre-certification measurements for five control-wafers from crystal 21566 vs. cumulative measurement run number.

Plot symbol code: 281 = 0; 2062 = 1; 2362 = 2; 283 = +, SRM1 = *

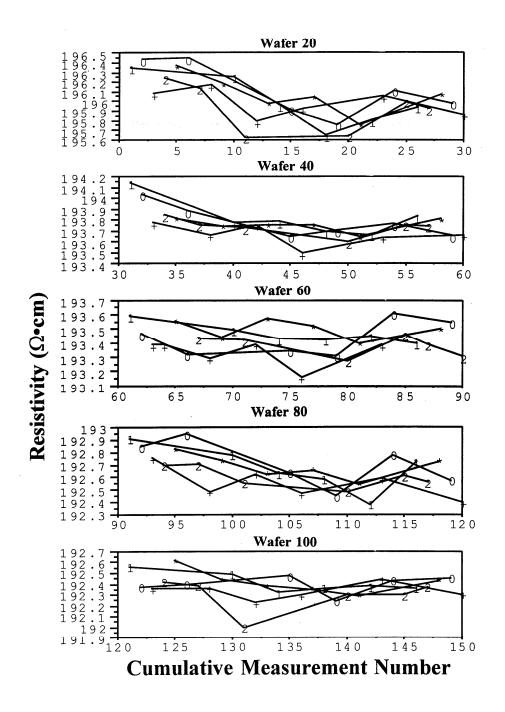


Figure 7. Resistivity (Ω ·cm) from post-certification measurements for five control-wafers from crystal 21566 vs. cumulative measurement run number.

Plot symbol code: 281 = 0; 2062 = 1; 2362 = 2; 283 = +, SRM1 = *

Reference:

Graybill, F. A., An Introduction to Linear Statistical Models, Vol. 1 (McGraw-Hill, New York, 1961), pp. 349-351.

Appendix 3. Analysis of Certification Data and Control Experiments for SRM 2541

1. GENERAL COMMENTS AND SUMMARY OF TYPE A STANDARD UNCERTAINTY COMPONENTS

1.1 Introduction

This appendix documents the statistical analysis leading to the certification of wafers from crystal 91905 for SRM 2541. The 130 wafers in this issue have nominal resistivities of $0.01~\Omega\cdot\text{cm}$, and the wafers are assumed to be identical with regard to face. For this issue, the pre- and post-certification measurements were made on opposite faces of each wafer. Certification measurements were made with probe 283.

This appendix includes a summary of the temporal and other components of uncertainty in Section 1.2, and details of the analysis for a systematic bias for probe 283 in Section 2.1. Such a probe-bias calculation was not illustrated in Appendix 2. The analyses of all other effects follow the procedures detailed in Appendix 2. The details are not included here.

1.2 Certified Resistivities and Uncertainties

The averages of six measurements on the 0 mm circle, and individual measurements on the 5 mm and 10 mm circles of each wafer are reported as certified values. No correction is applied for probe or wiring effects.

Only Type A uncertainty evaluation procedures are treated in this appendix, and estimates of all uncertainty components are shown in Table 1. The Type A standard uncertainty for the average resistivity at the wafer center is

$$u_i = \sqrt{\frac{b^2}{3} + s_{\gamma}^2 + s_{\delta}^2 + \frac{1}{6}s_{\varepsilon}^2} = 0.000\ 004\ 2\ \Omega \cdot \text{cm}\ (0.0042\ \text{m}\Omega \cdot \text{cm})\ .$$

Table 1. Components of Type A Standard Uncertainty for Crystal 91905, SRM 2541 with Probe 283, Q-cm

Type	Source	Std dev	Estimate
Random	Imprecision of probe 283	Se	0.000 001 83
Random	Run-to-run measurement variability	S_{δ}	0.000 001 04
Random	Long-term measurement variability	δ	0.000 004 00
Systematic	Uncertainty of zero correction for probe 283 bias	b/√3	0.000 000 47

Table 2. Sources of Variation for Crystal 91905, SRM 2541 with Probe 283, Ω·cm

Source of error	Experiment	RMSE a	DF^b	Relationship
	SRM certifications	0.000 001 70	650	
Probe imprecision	Pre-certification	0.000 002 30	150	, S
	Post- certification	0.000 002 02	150	
	Check standard	0.000 001 54	105	
	Pooled	0.000 001 83	1055	
	Pre- certification	0.000 001 59	25	
Run-to-run	Post-certification	0.000 001 20	25	$\binom{2}{S_c} + \frac{1}{Z} \binom{2}{S_c}$
	Check standard	0.000 000 87	20	(3,0,0,)
	Pooled	0.000 001 28	70	
Long-term	Pre- and post-certification	0.000 004 03	5	$(s_{\gamma}^{2} + \frac{1}{6}s_{\delta}^{2} + \frac{1}{36}s_{\varepsilon}^{2})^{1/2}$

associated with the individual effects, namely, imprecision, run-to-run variability, probe variability, and long-term variability, are computed ^aThe root-mean-square error, RMSE, estimates within each source-of-error category are pooled in the table above. Standard deviations using the relationship shown in the last column, with the results summarized in Table 1 of this. Appendix. ^bDegrees of Freedom.

^cThis column expresses the error components that comprise the pooled value in each "source of error" series of experiments; see reference at the end of Appendix 2. The Type A standard uncertainty for individual resistivity values on the 5 mm and 10 mm radius circles is

$$u_i = \sqrt{\frac{b^2}{3} + s_{\gamma}^2 + s_{\delta}^2 + s_{\epsilon}^2} = 0.000\ 004\ 5\ \Omega \cdot \text{cm}\ (0.004\ 5\ \text{m}\Omega \cdot \text{cm})$$
.

2. SYSTEMATIC EFFECTS

2.1 Bias Effect of Probe 283

There is a small systematic bias for this probe (relative to the average over all probes); the average bias is -0.000 000 68 Ω -cm with a standard deviation of the average of 0.0 000 18 Ω -cm. This bias can be seen in the measurements on the control-wafers, but does not affect the values of the SRMs which are only reported to six places beyond the decimal point. Therefore, the correction is taken to be zero. A conservative assumption is that during the certification the bias could fall somewhere within the limits $\pm b$ where 0.1 $b = 0.000 \ 000 \ 82 \ \Omega$ -cm, and a contribution of $b/\sqrt{3} = 0.000 \ 000 \ 47 \ \Omega$ -cm is included as a systematic component of the Type A standard uncertainty.

Table 3. Bias of Probe 283 Relative to the Average of All Probes, Ω·cm

Wafer	Pre-certification	Post-certification
2	-0.000 000 16	-0.000 001 32
43	-0.000 000 58	-0.000 000 58
44	0.000 000 48	-0.000 001 10
53	-0.000 001 44	-0.000 000 86
144	-0.000 000 94	-0.000 000 26
Mean	-0.000.000.53	-0 000 000 82

Appendix 4. Analysis of Certification Data and Control Experiments for SRM 2542

1. GENERAL COMMENTS AND SUMMARY OF TYPE A STANDARD UNCERTAINTY COMPONENTS

1.1 Introduction

This appendix documents the statistical analysis leading to the certification of wafers from crystal 91904 for SRM 2542. The 129 wafers in this issue have nominal resistivities of 0.1 Ω · cm, and the wafers are assumed to be identical with regard to face. For this issue, the pre- and post-certification measurements were made on opposite faces of each wafer. Certification measurements were made with probe 281.

This appendix includes a summary of the temporal and other components of uncertainty in Section 1.2, as well as the details of analysis of a term due to differences in probe-wiring configurations in Section 2.1. Such a probe-wiring calculation was not illustrated in Appendices 2 or 3. All other analyses follow procedures detailed in preceding appendices.

1.2 Certified Resistivities and Uncertainties

The averages of six measurements on the 0 mm circle, and individual measurements on the 5 mm and 10 mm circles of each wafer, corrected for wiring-configuration bias, are reported as certified values. No correction is applied for probe effect.

Only Type A uncertainty evaluation procedures are treated in this appendix, and estimates of all uncertainty components are shown in Table 1. The Type A standard uncertainty for the average resistivity at the wafer center is

$$u_i = \sqrt{\frac{b^2}{3} + s_{cfig}^2 + s_{\gamma}^2 + s_{\delta}^2 + \frac{1}{6}s_{\varepsilon}^2} = 0.000 \text{ 045 } \Omega \cdot \text{cm}.$$

The Type A standard uncertainty for the individual resistivity values on the 5 mm and 10 mm radius circles is

$$u_i = \sqrt{\frac{b^2}{3} + s_{cfig}^2 + s_{\gamma}^2 + s_{\delta}^2 + s_{\varepsilon}^2} = 0.000 \ 0.00$$

Table 1. Components of Type A Standard Uncertainty for Crystal 91904, SRM 2542 with Probe 281, Q.cm

Type	Source	Std dev	Estimate
Random	Imprecision of probe 281	Se	0.000 062
Random	Run-to-run measurement variability	$S_{\mathbf{\delta}}$	0.000032
Random	Long-term measurement variability	S_{γ}	0.000 004
Random	Standard deviation of correction for wiring configurations* s_{cfig}	s_{cfig}	0.000 011
Systematic	Uncertainty of a zero correction for probe 281 bias	p/\2	0.000 016

*A correction of -0.000 037 5 \Omega common due to wiring-configuration bias is applied to wafer center resistivity data for all wafers.

Table 2. Sources of Variation for Crystal 91904, SRM 2542 with Probe 281, Ω·cm

Source of error	Experiment	RMSE ^a	DF ^b	Relationship ^e
	SRM certifications	0.000 046	645	
Probe imprecision	Pre-certification	0.000 081	150	$\frac{S}{S}$
1	Post- certification	0.000 074	150	
	Check standard	980 0000	120	
	Pooled	0.000 062	1065	
	Pre- certification	0.000 028 9	15	2 1 2 1/2
Run-to-run	Post-certification	0.000 042 5	25	$(s^2 + s^2)$
	Check standard	0.000 044 6	23	
	Pooled	0.000 040 5	63	
Long-term	Pre- and post-certification	0.000 017	5	$\left(\frac{2}{s_{\gamma}} + \frac{1}{6}\frac{2}{s_{\delta}} + \frac{1}{36}\frac{2}{s_{\epsilon}}\right)^{1/2}$
	Pooled	0.000 017	5	

associated with the individual effects, namely, imprecision, run-to-run variability, probe variability, and long-term variability, are computed aThe root-mean-square error, RMSE, estimates within each source-of-error category are pooled in the table above. Standard deviations using the relationship shown in the last column, with the results summarized in Table 1 of this Appendix.

^bDegree of Freedom.

cThis column expresses the error components that comprise the pooled value in each "source of error" series of experiments; see reference at the end of Appendix 2.

2. SYSTEMATIC EFFECTS

2.1 Differences between Wiring Configurations b1 and b2

Differences are found between measurements in configurations b1 and b2. Averages and standard deviations (for the first four days of measurements on each wafer) are shown in Table 3. Rounds 5 and 6 of the pre-certification measurements were found to have been adversely affected by a faulty power supply that was discovered and repaired shortly after the start of wafer certification. Rounds 5 and 6 are omitted from the analysis of the probe-wiring effect for both pre- and post-certification control-wafer data. The t-statistic for testing for a significant difference between wiring configurations b1 and b2 is $t = \sqrt{20}$ Avg/SD. The values of the t-statistic suggest a slight difference between wiring configurations for this SRM. The average difference between the pre- and post-certification measurements is 0.000 075 Ω ·cm. A correction of minus one-half this difference, or -0.000 037 5 Ω ·cm, is applied to all certification measurements to obtain an average over the two configurations. The standard deviation of the correction,

$$s_{cfig} = \frac{1}{2} \frac{1}{\sqrt{20}} \sqrt{s_1^2 + s_2^2} = 0.000 \ 011 \ \Omega cm$$

where s_1 is the standard deviation from the pre-certification and s_2 is the standard deviation from post-certification measurement, is taken as a component of the Type A standard uncertainty for the process.

Table 3. Average Differences and Standard Deviations between Wiring Configurations b1 and b2, Ω•cm

	Pre-c	ertification			Post-certification			
Probe	Avg	$SD(s_1)$	DF	t	Avg	$SD(s_2)$	DF	t
281	0.000 085	0.000 064	19	5.9	0.000 065	0.000 072	19	4.0

Appendix 5. Analysis of Certification Data and Control Experiments for SRM 2545

1. GENERAL COMMENTS AND SUMMARY OF TYPE A STANDARD UNCERTAINTY COMPONENTS

1.1 Introduction

This appendix documents the statistical analysis leading to the certification of wafers from crystal 21565, SRM 2545. The 133 wafers in this issue have nominal resistivities of $25 \,\Omega$ cm, and the wafers are assumed to be identical with regard to wafer face. For this SRM, the pre- and post-certification measurements were made on opposite faces of each wafer. Certification measurements were made with probe 2062.

This appendix includes a summary of the temporal and other components of uncertainty in Section 1.2, as well as details of an analysis for wiring-configuration differences of a form not contained in any of the previous appendices. All other analyses follow procedures detailed in Appendix 2.

1.2 Certified Resistivities and Uncertainties

The averages of six measurements on the 0 mm circle, and individual measurements on the 5 mm and 10 mm circles of each wafer, are reported as certified values. There is no correction for probe effect.

Only Type A uncertainty evaluation procedures are treated in this appendix, and estimates of all uncertainty components are shown in Table 1. The Type A standard uncertainty for the average resistivity at the wafer center is

$$u_i = \left(\frac{a^2}{3} + s_{\gamma}^2 + s_{\delta}^2 + \frac{1}{6}s_{\varepsilon}^2\right)^{1/2} = 0.008 \ \Omega \cdot \text{cm} .$$

The Type A standard uncertainty for the individual resistivity values on the 5 mm and 10 mm radius circles is

$$u_i = \left(\frac{a^2}{3} + s_{\gamma}^2 + s_{\delta}^2 + s_{\varepsilon}^2\right)^{1/2} = 0.015 \,\Omega \cdot \text{cm}.$$

Table 1. Components of Type A Standard Uncertainty for Crystal 21565, SRM 2545 with Probe 2062, Ω·cm

Type	Source	Std dev	Std dev Estimate
Random	Imprecision of probe 2062	Sc	0.014 14
Random	Run-to-run measurement variability	s s	0.003 31
Random	Long-term measurement variability	. S.	0.003 01
		-	
Systematic	Uncertainty of a zero correction for differences between wiring configurations a//3	a/√3	0.002 89

Table 2. Sources of Variation for Crystal 21565, SRM 2545 with Probe 2062, Ω·cm

Source of error	Experiment	RMSE ^a	DF	Relationship ^c
	SRM certifications	0.014 93	059	
Probe imprecision	Pre-certification	0.011 32	150	S
	Post-certification	0.014 67	150	
	Check standard	0.012 04	120	
	Pooled	0.014 14	1070	
	Pre-certification	896 900.0	25	
Run-to-run	Post-certification	0.004 588	25	$(s_{\delta}^2 + \frac{1}{5}s_{\epsilon}^2)$
	Check standard	0.008 056	23	,
	,			
	Pooled	0.006 655	73	
Long-term	Pre- and post-certification	0.004 057	$\frac{2}{s}$	$(s_{\gamma}^{2} + \frac{1}{6}s_{\delta}^{2} + \frac{1}{36}s_{\epsilon}^{2})^{1/2}$
	Pooled	0.004 057	5	

associated with the individual effects, namely, imprecision, run-to-run variability, probe variability, and long-term variability, are computed from ^aThe root-mean-square error, RMSE, estimates within each source-of-error category are pooled in the table above. Standard deviations the relationships shown in the last column, with the results summarized in Table 1 of this Appendix.

^bDegrees of Freedom.

^cThis column expresses the error components that comprise the pooled value in each "source of error" series of experiments; see reference at the end of Appendix 2.

2. SYSTEMATIC EFFECTS

2.1 Differences between Wiring Configurations b1 and b2

Differences are found between measurements in configurations b1 and b2. An obvious outlier in the pre-certification measurements on wafer 39 was deleted from the database for the purpose of the analysis. Averages and standard deviations are shown in Table 3. The t-statistic for testing for a significant difference between wiring configurations b1 and b2 is $t = \sqrt{29}$ Avg/SD. The t-statistics suggest a slight difference among wiring configurations for this issue, although the differences are in opposite directions for the pre- and post-certification measurements. With no other information at hand, it is reasonable to assume that during the certification procedure, the difference between wiring configurations could fall somewhere within the limits $\pm a$, where $a = 0.005 \ \Omega \cdot cm$ is based on the post-certification average value. It is also reasonable to assume that the best correction is zero, and that the standard uncertainty for the underlying uniform distribution is $a/\sqrt{3}$, or $0.002 \ 89 \ \Omega \cdot cm$.

Table 3. Average Differences and Standard Deviations between Wiring Configurations b1 and b2, Ω•cm

	Pre-c	certification			Pe	Post-certification			
Probe	Avg	SD	DF	t	Avg	SD	DF	t	
2062	-0.003 83	0.005 14	28	-4.0	+ 0.004 89	0.004 00	28	6.6	

Appendix 6. Analysis of Certification Data and Control Experiments for SRM 2546

1. GENERAL COMMENTS AND SUMMARY OF TYPE A STANDARD UNCERTAINTY COMPONENTS

1.1 Introduction

This appendix documents the statistical analysis leading to the certification of wafers from crystal 51939 for SRM 2546. The 130 wafers in this issue have nominal resistivities of 100 Ω ·cm, and the wafers are assumed to be identical with regard to face; all measurements were made on the same face of each wafer. All certification measurements were made with probe 2362.

This appendix contains a summary of the temporal and other components of uncertainty in Section 1.2. All analyses of the temporal components of uncertainty for this SRM follow procedures detailed in Appendix 2 for analysis of SRM 2547. Section 2.1 summarizes an analysis of a probe bias correction that follows the procedures used in Appendix 2.

1.2 Certified Resistivities and Uncertainties

The averages of six measurements on the 0 mm circle, and individual measurements on the 5 mm and 10 mm circles of each wafer, corrected for probe #2362 are reported as certified values.

Only Type A uncertainty evaluation procedures are treated in this appendix, and estimates of all uncertainty components are shown in Table 1. The Type A standard uncertainty for the average resistivity at the wafer center is

$$u_i = \left(s_c^2 + s_{\gamma}^2 + s_{\delta}^2 + \frac{1}{6}s_{\varepsilon}^2\right)^{1/2} = 0.036 \,\Omega \cdot \text{cm}.$$

The Type A standard uncertainty for individual resistivity values on the 5 mm and 10 mm radius circles is

$$u_i = (s_c^2 + s_v^2 + s_\delta^2 + s_\varepsilon^2)^{1/2} = 0.075 \text{ cm}.$$

Table 1. Components of Type A Standard Uncertainty for Wafers from Crystal 51939, SRM 2546 with Probe 2362, Ω·cm

Type	Source	Std dev	Estimate
Random	Imprecision of probe 2362	$S_{\mathbf{E}}$	0.0720
Random	Run-to-run measurement variability	S_{δ}	0.0134
Random	Long-term measurement variability	S_{γ}	0.0146
Random	Standard deviation of the correction for probe 2362 bias*	S_C	0.0051

*A correction of $0.0393\,\Omega$ cm due to probe 2362 bias is applied wafer-center resistivity data for all wafers.

Table 2. Sources of Variation for Wafers from Crystal 51939, SRM 2546 with Probe 2362, Q.cm

ומטוע ליי	table 2. Doubles of Furtation for Walets month Ciferia 21/37, Start 2340 With 1100c 2302, St VIII	total City state 217	77, DIMI 27TO V	With 1 1000 2002, 32 Oill
Source of error	Experiment	RMSE ^a	DF^{b}	Relationship ^e
	SRM certifications	0.074 6	059	
Probe imprecision	Pre- and post-certification	0.071 0	300	38
	Check standard	0.0596	125	
	Pooled	0.072 0	1075	
Run-to-run	Pre- and post-certification	0.0362	50	$(s_{\rm s}^2 + \frac{1}{\epsilon} s_{\rm s}^2)^{1/2}$
	Check standard	0.0268	24	(3-0 0-)
	Pooled	0.032 3	74	
Long-term	Pre- and post-certification	0.0197	5	$\left(s_{\gamma}^{2}+\frac{1}{6}s_{\delta}^{2}+\frac{1}{36}s_{\epsilon}^{2}\right)^{1/2}$
	Pooled	0.019 7	5	

associated with the individual effects, namely, imprecision, run-to-run variability, probe variability, and long-term variability, are computed from ^aThe root-mean-square error, RMSE, estimates within each source-of-error category are pooled in the table above. Standard deviations the relationships shown in the last column, with the results summarized in Table 1 of this Appendix.

bDegrees of Freedom.

^cThis column expresses the error components that comprise the pooled value in each "source of error" series of experiments; see reference at the end of Appendix 2.

2. SYSTEMATIC EFFECTS

2.1 Bias Effect of Probe 2362

Differences from the multi-probe mean were found for probe 2362 for each wafer, and are given in Table 3. The estimated correction for this probe over five wafers is $\hat{\mathbf{C}} = +0.0393~\Omega\cdot\mathrm{cm}$; the standard deviation of this average correction is $s_c = 0.0051~\Omega\cdot\mathrm{cm}$. The correction, $\hat{\mathbf{C}}$, is applied to all certified values, and its standard deviation is taken as a component of the Type A standard uncertainty.

Table 3. Bias of Probe 2362 Relative to the Average for All Probes, Ω·cm

Wafer#	Pre-certification	Post-certification
138	0.0372	0.0507
139	0.0094	0.0657
140	0.0261	0.0398
141	0.0252	0.0534
142	0.0383	0.0469

Mean Bias, \hat{C} 0.0393 Ω·cm Standard Deviation of Mean 0.0051 Ω·cm

Appendix 7. SRM Values after an Extended Period of Time

It is useful to evaluate how closely the original SRM measurement scale can be reproduced after extended periods of time since the certification and control measurements were originally taken. To do this, a single set of six measurements was taken at the center of each of the original control wafers from the pre- and post-certification experiments with the same probe used originally. None of the probes had been rebuilt or modified since the earliest of the certification measurements, those at 200 Ω cm, for SRM 2547. However, all probes had been used at least for all of the pre- and post-certification control experiments, and probe 283 was used for all certification measurements for three of the SRM levels including the very first and the very last ones measured. As a result, some wear can be expected on all probes between the time they were first used and the time of the recent follow-up measurements.

1. Summary of Results

The results are summarized in Table 1 where the entries in the second and fourth columns are the grand averages from six runs using the probe noted during the pre- or post-certification control experiments. The entries in the third and fifth columns are the standard deviations, Σ , of the average values from each of those six runs. The value in column six is the average of six measurements at the wafer center during a single run in February 1997. The entry in the seventh column is the standard deviation, σ , of those six individual measurements. Finally, the value in the last column is the relative difference between the single average value from February 1997 and the grand average, or base-line value, of all 12 runs from the pre- and post-measurement experiments. Data from pre- and post-measurement experiments are stated as actually acquired. No correction for probe bias or for configuration bias, such as are identified in some of the statistical analysis reports as being necessary for the SRM wafers to be issued, was applied.

2. Comments on the Results

For measurements on wafers at 0.01 Ω cm, 0.1 Ω cm, 10 Ω cm, 25 Ω cm, and 100 Ω cm, the latest measurements appear to be randomly above and below the base-line values from the control experiments. With the exception of wafer #141 at 25 Ω cm, recent measurements at those resistivities are all within 0.10 % of the base-line values. Measurements on wafers at 1 Ω cm show a consistent high-side bias of recent values over the base-line results. Because of the known residual sensitivity of the 1 Ω cm material to illumination levels, it must be considered that a difference in illumination levels between that during the latest measurements and that at the time of the original measurements could be responsible for causing all measurement differences to be of the same sign. Measurements at 200 Ω cm also show a systematic difference between recent and base-line values. In this case, present values are below the base-line values, in the direction of the shift with remeasurement previously noted. While this effect may be the dominant cause of the observed shift, average relative humidity was approximately 45 % at the time of the base-line measurements and was approximately 32 % during the latest measurements. In order to put the latest values in perspective, an additional column has been added to the table for the 200 Ω cm wafers. This column gives

the two standard deviation (2 σ) lower limit value for each wafer, which is calculated from the base-line average value and the lower 2 σ uncertainty value given in Appendix 2. It can be seen that the February 1997 values are clearly within the lower 2 σ limit for the 200 Ω cm SRM level.

Table 1. Summary of the Six-Round Grand Averages and Standard Deviations from Preand Post-Certification Measurements, Single Round Averages and Standard Deviations from Recent Measurements, and the Percent Changes in Measurement Values

CRYSTAL 91905 Probe 283 Elapsed Time 38 Months

							Feb 1997
Control	Pre-certification		Post-certifi	cation	Feb. 1997		Minus
Wafer#	· avg		$ ho_{avg}$		$ ho_{ m avg}$		Pre/Post Avg
	$(\Omega \text{ cm})$	$(\Sigma,\%)$	$(\Omega^{c} cm)$	$(\Sigma,\%)$	$(\Omega \cdot \mathbf{cm})$	(σ,%)	(Difference, %)
002	0.011 286	0.021	0.011 275	0.015	0.011 281	0.046	+0.004
043	0.010 974	0.013	0.010 972	0.014	0.010 970	0.024	-0.027
044	0.010 955	0.010	0.010 949	0.004	0.010 954	0.030	+0.018
053	0.010 923	0.015	0.010 926	0.006	0.010 923	0.023	-0.014
144	0.010 350	0.014	0.010 352	0.012	0.010 352	0.015	+0.010
2	0.010 350	0.011	0.010 302	0.012	0.010 302	0.015	. 0.010
•		CRYSTAL	01004 P*	obe 281	Flanced Tim	10 Mo	
		CKISIAL	7 7 1 7 U4 F F	ODE 201	Elapsed Tim	ie 19 Mioi	IIIIS
003	0.114 51	0.137	0.114 59	0.039	0.114 59	0.091	+0.035
066	0.113 88	0.123	0.113 82	0.063	0.113 80	0.110	-0.044
097	0.112 61	0.143	0.112 52	0.016	0.112 56	0.062	-0.004
161	0.104 35	0.146	0.104 33	0.012	0.104 32	0.076	-0.019
287	0.099 61	0.129	0.099 60	0.035	0.099 63	0.047	+0.025
-			<u> </u>				
		CRYSTAL	. 91907 Pr	obe 283	Elapsed Tim	ie 31 Moi	nths
011	1.0733	0.028	1.0733	0.059	1.0745	0.068	+0.110
026	1.0605	0.023	1.0607	0.024	1.0613	0.022	+0.066
042	1.0461	0.035	1.0463	0.014	1.0470	0.078	+0.076
131	0.9911	0.033	0.9916	0.014	0.9916	0.077	+0.025
208	0.9619	0.033	0.9623	0.025	0.9630	0.125	+0.094

Table 1 (cont'd.)

CRYSTAL 29473 Probe 283 Elapsed Time 3 Months

	ol Pre-certi	fication	Post-c	ertification	Feb 1997	7	Feb 1997 Minus
Wafe	r# ρ _{avg} (Ω·cm)	(Σ,%)	ρ _{avg} (Ω cn	n) $(\Sigma,\%)$	$ ho_{ m avg} \ (\Omega{ m cm})$	(σ,%)	Pre/Post Avg (Difference, %)
016	10.085	0.014	10.080	0.033	10.085	0.079	+0.025
032	10.105	0.025	10.096	0.021	10.109	0.049	+0.084
075	10.316	0.029	10.308	0.024	10.309	0.085	-0.029
108	10.186	0.017	10.177	0.020	10.181	0.041	-0.005
120	10.082	0.017	10.073	0.014	10.077	0.056	-0.005
	-			·			· · · · · · · · · · · · · · · · · · ·
		CRYSTA	L 21565	Probe 2062	Elapsed Ti	ime 29 N	Tonths
017	24.050	0.032	24.046	0.015	24.046	0.104	-0.008
039	24.695	0.029	24.699	0.022	24.701	0.061	+0.016
063	24.509	0.016	24.517	0.011	24.495	0.058	-0.073
103	24.135	0.031	24.142	0.025	24.124	0.044	-0.060
125	24.052	0.032	24.056	0.019	24.054	0.068	+0.001
			-		-		
		CRYSTA	L 51939	Probe 2362	Elapsed T	ime 34 N	Months
138	95.093	0.038	95.124	0.048	95.131	0.125	+0.024
139	99.306	0.048	99.310	0.022	99.252	0.076	-0.056
140	96.036	0.028	96.077	0.029	96.103	0.072	+0.048
141	101.060	0.023	101.079	0.053	101.277	0.097	+0.205
142	94.215	0.029	94.244	0.039	94.309	0.080	+0.084

CRYSTAL 21566 Probe SRM1 Elapsed Time 54 Months

Control	Pre-certification		Post-certification		Feb 1997		Feb 1997 Minus	2σ
Wafer#	$ \rho_{avg} $ (Ωcm)	(Σ,%)	$ \rho_{avg} $ $(\Omega \cdot cm)$	$(\Sigma,\%)$	$_{(\Omega\text{ cm})}^{\rho_{avg}}$	(σ,%)	Pre/Post Avg (Difference, %)	Lower Limit
020	196.27	0.050	196.07	0.104	196.05	0.081	-0.061	195.60
040	193.88	0.034	193.76	0.032	193.59	0.102	-0.119	193.39
060	193.57	0.072	193.50	0.036	193.24	0.039	-0.152	193.10
080	192.82	0.054	192.69	0.050	192.35	0.097	-0.210	192.32
100	192.59	0.065	192.42	0.057	192.26	0.074	-0.127	192.07